

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 14:46:57 ; Search time 7316.32 Seconds

(without alignments)
4710.839 Million cell updates/sec

Title: US-09-826-581-5

Perfect score: 1647

Sequence: 1 ttggtctggggctggccaca.....acacacagctctagctcttc 1647

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : GenBank.*
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3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
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10: gb_ro.*
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12: gb_un.*
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14: gb_vl.*
15: em_ba.*
16: em_fun.*
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18: em_in.*
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22: em_ov.*
23: em_pat.*
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25: em_pl.*
26: em_ro.*
27: em_sts.*
28: em_un.*
29: em_vl.*
30: em_htg_hum.*
31: em_htg_inv.*
32: em_htg_other.*
33: em_htgo_inv.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result Query #
No. Score Match Length DB ID Description

1	1647	100.0	1647	6	AX281582	AX281582 Sequence
2	1590	96.5	2290	6	HS2429977	HS2429977 Homo sapi
3	1453	88.2	2115	6	AX099802	AX099802 Sequence
4	1453	88.2	2115	9	AF214519	AF214519 Homo sapi
5	1447	87.9	2109	6	AX099776	AX099776 Sequence
6	1172.8	71.2	2022	6	AX099804	AX099804 Sequence
7	1140.4	69.2	1873	4	AF214520	AF214520 Sus scrofa
8	1140.4	69.2	1873	6	AX099800	AX099800 Sequence
9	1134.4	68.9	1867	6	AX099774	AX099774 Sequence
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11	414.8	25.2	1328	10	RNMPKGM	X95578 R. norvegicus
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14	396.8	24.1	1623	10	AF035635	AF035635 Mus muscu
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17	393.4	23.9	1578	9	HS042413	U42412 Human 5'-AM
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19	366.6	22.3	1167	9	AB025580	AB025580 Homo sapi
20	366.6	22.3	1435	6	AR139104	AR139104 Sequence
21	366.6	22.3	2194	9	AF087875	AF087875 Homo sapi
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24	363.4	22.1	2062	9	HS2429976	AF094764 Drosophila
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27	278.8	16.9	1014	6	AC027416	AC027416 Homo sapi
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ALIGNMENTS

RESULT 1
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LOCUS AX281582 1647 bp DNA linear PAT 02-NOV-2001
DEFINITION Sequence 5 from Patent WO0177305.
ACCESSION AX281582
VERSION AX281582.1 GI:16608833
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
1 (sites)
Andersson, L., Luthman, H. and Marklund, S.
TITLE Variants of the human amp activated protein kinase gamma 3 subunit
JOURNAL Patent: WO 0177305-A 5 18-Oct-2001;
Atexis AB (SE)
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CDS

file
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Sequence search results

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Query Match	100.0%;	Score 1647;	DB 6;	Length 1647;
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Matches 1647; Conservative	0;	Mismatches	0;	Indels 0; Gaps 0;

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QY	361	ggactgcctccctctgactgagtacagcctcaagctgcagaggttccagaacagatgagtgtga	42
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QY	421	gctggcccaaggaggttcccagccacaagagcctctggagttgtgagctagaagaagcctctga	48
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TITLE

Characterization of AMP-activated protein kinase gamma-subunit isoforms and their role in AMP binding

JOURNAL

Biochem. J. 346 Pt 3, 659-669 (2000)

MEDLINE

20164049

REFERENCE

2 (bases 1 to 2290)

AUTHORS

Carling, D.

JOURNAL

Submitted (12-OCT-1999) Carling D., Cellular Stress Group, MRC Clinical Sciences Centre, Hammersmith Hospital, DuCane Road, London, W12 0NN, UNITED KINGDOM

FEATURES

source

1. .2290

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22. .1500

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LSPOAFPFKIGMDDELKPKGAOYIMREIEHCTCDYAMATSSVLEFDPMLETKKAPFA

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BASE COUNT

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ORIGIN

Query Match

Best Local Similarity 96.5%; Score 1590; DB 9; Length 2290;

Matches 1607; Conservative 0; Mismatches 10; Indels 1; Gaps 1;

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RESULT 3
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DEFINITION Sequence 29 from Patent WO0120003.
ACCESSION AX099802
VERSION AX099802.1 GI:13538836
KEYWORDS human.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
Andersson, L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Rogel-Gallard, C., Iannucci, N., Gellin, J., Le Roy, P., and
Chardon, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 29 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
Location/Qualifiers
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BASE COUNT 460 a 622 c 562 g 471 t
ORIGIN

Query Match 88.2%; Score 1453; DB 6; Length 2115;
Best Local Similarity 97.3%; Pred. No. 0;
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215 gtgaggaagggagccacaggttcagggggaaggtcccggttcagagccagcttgtag 274
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335 ggggttgagcactccacacaggggtgagatctccctccttgatctgtagacagcccaact 394
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395 ggaagctcagacagatgtagtgagctggccaggaagttccagcacagagggcctgg 454

301 GCAGGCTCCAGCAGACATGATGTGAGAGCTGGCCAGGAGTTCCACAGAGGCGCTGG 360
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DEFINITION	Sequence	from Patent WO0120003.	
VERSION	AX099802		
KEYWORDS	AX099802.1	GI:13538836	
SOURCE	.	human.	
ORGANISM	human sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
TITLE	1 (bases 1 to 2115)		
JOURNAL	Anderson, L., Looft, C., Kaim, E., Milan, D., Robic, A., Rogel-Gallard, C., Iannucci, N., Gellin, J., Le Roy, P. and Chardon, P. Variants of the gamma chain of ampk, dna sequences encoding the same, and uses thereof Patent: WO 0120003-A 29 22-MAR-2001; INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ; Andersson, Leif (SE) ; Looft, Christian (DE) ; Kaim, Ernst (DE) Location/Qualifiers 1..2115 /organism="Homo sapiens" /db_xref="taxon:9606" 1..1395 /note="unnamed protein product" /codon_start=1 /protein_id="CAC35801.1" /db_xref="GI:13538837" /translation="MSFLEQENSSSWSPATVSSSRIRGKRAKLFWTRFKSVEEG EPPOGEPRPRTPESTGEATFPKPTPLAQADPADGTPTGMCDLPSCTSAAS SSTDVELATEFPATEAMECELEGLERPALCLSPAPRPKLMDRLRGAOIYV RFMEHCYDAMATRSKLVIFDTMLEIKAFALVANCPRAAPLMDSKOSFGMLT TDFLVLHRYRSLVDLYELEIOHKIEWREIYLGGCKPLVISPNDSLFEAYTLT KNRIHRLPVLDVPVSGNVLHTHRKLKLFHI FGSILPRPSFLTIDDLIGTFROI AVLETAPALITDALDPDRVSALPVNEGGQWGLYSRPVTHLAODTYNHLDMS GEALRORTICLEGDILOSCPHRESLDEVDIRAREOYHRLVLYDETQHLLGVLSLDILA ALVSPAGIDMLGA"		
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Best Local Similarity	97.3%; Pred. No. 0;		
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Dd	121	GTGAGAGAAAGGGAGGCCACCAGGTCAGGGGCAAGGTCCC CGTCCAGGCCAACTGCTAG	180
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Dd	181	TCACACGGGGCTGGAGGCCACATTCCCAAAGCCACACCTTTGGCTCAAGCGATCTCTCC	240
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Db	361	GAGGTGAGACTGAAAGGCTCTGCTGGAAAGAGAGGCTTGCCCTGTGCTGTCTCCCGCAGGCC	420
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Qy	575	gcctctatgcagagagcaaacctgtctaaatgacatggcaatggcaactatgcttcaagctaatgac	634
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Qy	755	ttcacctccggtgctgacatcagctactaagatgccccctgtgtccagatctatgagatctgaa	814
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Qy	815	caacaataagattgagaccctggaagagagatcactctgaagcctgtcttcaagcctctgtgtc	874
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Qy	1055	cgcactatccaaagatttgagcatgagacatccgaagacttgctgtgtgtctgtaga	1115
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Db	1021	GCACCCATCCGATGTGCACATGTCGATCTTTGTGTGGACCGGGCTGTCTCTCACTGCTGTG	1080
Qy	1175	gtcaacgaatgtgttcaagatgcttgagggcctcatctccgagcttgaatgtatcaactgagct	1235
Db	1081	GTCAACGAATGTGGTGCAGGTGTGTGGGCTCTATTCCCGCTTTGATGTGATTCACTGGCT	1140
Qy	1235	gcceagcaaacctacaacacacttgagacaatgagtgtggaagaagcccttgagcagagaca	1295
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Qy	1295	ctatgtctgagaggaagtccttctctgcgaagccccaagaagacttgagggaatgtatgac	1355
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Qy	1355	aggaatgctcggagacagatcacacagagctgtgtctagtgtagagagaccgaagatctctg	1415
Db	1261	AGGATTTCTCGGAGCGAGTACACAGAGCTGTGCTAGTGAAGAGACAGACATCTTTTG	1320
Qy	1415	ggagtggtctcccccctccgaacatccttctgaagcctgtgtctgaagccctgtgtgactgat	1475
Db	1321	GGCTGTGCTCTCTCTCCGACATCTTCAAGGCACTGTGCTCTCAAGCCCTGTGCGCATTCAT	1380
Qy	1475	ggccctcggggcttgagaagaatctcgaatctccatcatcccaagcaccctgcacacacttgagac	1535
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LOCUS	2115 bp mRNA linear PRI 03-JUN-2000
DEFINITION	Homo sapiens AMP-activated protein kinase gamma subunit (PRKAG3)
ACCESSION	AF214519
VERSION	AF214519.1 GI:8215681
KEYWORDS	human.
SOURCE	

REFERENCE
AUTHORS
Milan, D., Jeon, J.-T., Looft, C., Amaral, V., Robic, A., Thelander, M.,
Mammalla, Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 2115)

TITLE	A mutation in PRKAG3 associated with excess glycogen content in pig skeletal muscle
JOURNAL	Science 288 (5469), 1248-1251 (2000)
PMID	20280150
PUBMED	10818001

AUTHORS Milán, D., Jeon, J.-T., Looft, C., Amarger, V., Robic, A., Rogel-gallard, C., Paul, S., Gellin, D., Lundström, K., Kalm, E., Le Roy, P., Chardon, P. and Andersson, L.

TITLE Direct Submission

JOURNAL Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish University of Agricultural Sciences, BMC box 597, Uppsala 751 24, Sweden

FEATURES	source	Location/Qualifiers
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Best Local Similarity	97.3%	Pred. No. 0		
Matches 1501; Conservative	0	Mismatches 35	Indels 7	Gaps 2

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QY	215	gttgaggaagagggaagccacacaggttcaaggggaaagtctcccggtctccaggccagctgtgag	274
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QY	1055	cgaactatcaagaatttgggcatcggcaacttcggagacttggcctgtgtgtctgtgagaca	1114
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LOCUS AX099776
DEFINITION Sequence 3 from Patent WO0120003.
ACCESSION AX099776
VERSION AX099776.1 GI:13538810
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 2109)
ANDERSSON, L., LOOF, C., KALM, F., MILAN, D., ROBIC, A.,
ROGEL-GALLIARD, C., IANNUCELLI, N., GELLIN, J., LE ROY, P. and
CHARDON, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 3 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) (DE)
ANDERSSON, Lelf (SE) ; LOOF, Christian (DE) ; KALM, Ernst (DE)
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BASE COUNT 458 a 621 c 560 g 470 t
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ORGANISM	Sus scrofa		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
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REFERENCE	1 (bases 1 to 2022)		
AUTHORS	Andersson, U., Looft, C., Kalm, E., Milan, D., Robic, A.,		
	Rogel-galliard, C., Iannuccelli, N., Gellin, J., Le Roy, P. and		
	Chardon, P.		
TITLE	Variaants of the gamma chain of ampx, dna sequences encoding the		
	same, and uses thereof		
JOURNAL	Patent: WO 0120003-A 31 22-MAR-2001;		
	INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;		
	Andersson, Lelf (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)		
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RESULT 7
AF214520

LOCUS 1873 bp mRNA linear MAM 03-JUN-2000

DEFINITION Sus scrofa AMP-activated protein kinase gamma subunit (PRKAG3)

ACCESSION AF214520

VERSION AF214520.1 GI:8215683

KEYWORDS pig.

SOURCE Sus scrofa

ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.

REFERENCE 1 (bases 1 to 1873)
Mylan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Thelander,M., Rogel-Gallard,C., Paul,S., Gellin,J., Lundstrom,K., Reinsch,N., Lundstrom,K., Reinsch,N., Gellin,J., Kalm,E., Roy,P.L., Chardon,P., and Andersson,L.
A mutation in PRKAG3 associated with excess glycogen content in pig skeletal muscle
Science 288 (5469), 1248-1251 (2000)

TITLE JOURNAL
MEDLINE 20280150
PUBMED 10818001
2 (bases 1 to 1873)
Mylan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Thelander,M., Rogel-Gallard,C., Paul,S., Gellin,J., Lundstrom,K., Reinsch,N., Kalm,E., Le Roy,P., Chardon,P., and Andersson,L.
Direct Submission
Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish University of Agricultural Sciences, BMC box 597, Uppsala 751 24, Sweden

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ACCESSION AX099774
VERSION AX099774.1 GI:13538808
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
1 (bases 1 to 1867)
Andersson, L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Rogel-Gaillard, C., Iannucci, J. N., Geil, J., Le Roy, P., and
Chardon, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof

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JOURNAL Patent: WO 0120003-A 1 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
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DEFINITION kinase.
ACCESSION X95578.1 GI:1185270
VERSION X95578
KEYWORDS AMP-activated protein kinase; gamma subunit.
SOURCE Norway rat.
ORGANISM Rattus norvegicus
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE 1 (bases 1 to 1328)
AUTHORS Woods,A., Cheung,P.C., Smith,F.C., Davison,M.D., Scott,J.,

TITLE Bert,R.K. and Carling,D.
JOURNAL Characterization of AMP-activated protein kinase beta and gamma
MEDLINE subunits: Assembly of the heterotrimeric complex in vitro
REFERENCE J. Biol. Chem. 271 (17), 10282-10290 (1996)
96215327
AUTHORS 2 (bases 1 to 1328)
TITLE Carling,D.
JOURNAL Direct Submission
Submitted (07-FEB-1996) D. Carling, MRC Clinical Sciences Centre,
Department of Molecular Medicine, RPMs, Hammersmith Hospital,
Ducane Road, London, W12 0NN, UK
FEATURES
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BASE COUNT 330 a 337 c 340 g 321 t
ORIGIN

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Matches 614; Conservative 0; Mismatches 332; Indels 0; Gaps 0;

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Db 301 GACTTATATCAATATTCTGACCGGATATCTCAAGTCAAGCCCTGTGTGACATCTATCACTG 360
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QY 872 gtctcactctctcctaataatagatagcctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 931
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REFERENCE		<p>1. (bases 1 to 1550) Gao, G., Fernandez, C.S., Stapleton, D., Auster, A.S., Widmer, J., Dyck, J.R., Kemp, B.E. and Witters, L.A. Non-catalytic beta- and gamma-subunit isoforms of the 5'-AMP-activated protein kinase J. Biol. Chem. 271 (15), 8675-8681 (1996)</p>
JOURNAL MEDLINE REFERENCE		<p>2 (bases 1 to 1550) Gao, G., Widmer, J., Auster, A., Stapleton, D.S., Kemp, B.E. and Witters, L.A. Direct Submission Submitted (07-DEC-1995) Lee A. Witters, Medicine/Biochemistry, Dartmouth Medical School, N. College St., Hanover, NH 03755-3833, USA</p>
TITLE		Location/Qualifiers
JOURNAL		1. 1350
REFERENCE		<p>1. (bases 1 to 1550) Gao, G., Fernandez, C.S., Stapleton, D., Auster, A.S., Widmer, J., Dyck, J.R., Kemp, B.E. and Witters, L.A. Non-catalytic beta- and gamma-subunit isoforms of the 5'-AMP-activated protein kinase J. Biol. Chem. 271 (15), 8675-8681 (1996)</p>
JOURNAL MEDLINE REFERENCE		<p>2 (bases 1 to 1550) Gao, G., Widmer, J., Auster, A., Stapleton, D.S., Kemp, B.E. and Witters, L.A. Direct Submission Submitted (07-DEC-1995) Lee A. Witters, Medicine/Biochemistry, Dartmouth Medical School, N. College St., Hanover, NH 03755-3833, USA</p>
TITLE		Location/Qualifiers
JOURNAL		1. 1350
REFERENCE		<p>1. (bases 1 to 1550) Gao, G., Fernandez, C.S., Stapleton, D., Auster, A.S., Widmer, J., Dyck, J.R., Kemp, B.E. and Witters, L.A. Non-catalytic beta- and gamma-subunit isoforms of the 5'-AMP-activated protein kinase J. Biol. Chem. 271 (15), 8675-8681 (1996)</p>
JOURNAL MEDLINE REFERENCE		<p>2 (bases 1 to 1550) Gao, G., Widmer, J., Auster, A., Stapleton, D.S., Kemp, B.E. and Witters, L.A. Direct Submission Submitted (07-DEC-1995) Lee A. Witters, Medicine/Biochemistry, Dartmouth Medical School, N. College St., Hanover, NH 03755-3833, USA</p>
TITLE		Location/Qualifiers
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JOURNAL MEDLINE REFERENCE		<p>2 (bases 1 to 1550) Gao, G., Widmer, J., Auster, A., Stapleton, D.S., Kemp, B.E. and Witters, L.A. Direct Submission Submitted (07-DEC-1995) Lee A. Witters, Medicine/Biochemistry, Dartmouth Medical School, N. College St., Hanover, NH 03755-3833, USA</p>
TITLE		Location/Qualifiers
JOURNAL		1. 1350
REFERENCE		<p>1. (bases 1 to 1550) Gao, G., Fernandez, C.S., Stapleton, D., Auster, A.S., Widmer, J., Dyck, J.R., Kemp, B.E. and Witters, L.A. Non-catalytic beta- and gamma-subunit isoforms of the 5'-AMP-activated protein kinase J. Biol. Chem. 271 (15), 8675-8681 (1996)</p>
JOURNAL MEDLINE REFERENCE		<p>2 (bases 1 to 1550) Gao, G., Widmer, J., Auster, A., Stapleton, D.S., Kemp, B.E. and Witters, L.A. Direct Submission Submitted (07-DEC-1995) Lee A. Witters, Medicine/Biochemistry, Dartmouth Medical School, N. College St., Hanover, NH 03755-3833, USA</p>
TITLE		Location/Qualifiers
JOURNAL		1. 1350
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JOURNAL MEDLINE REFERENCE		<p>2 (bases 1 to 1550) Gao, G., Widmer, J., Auster, A., Stapleton, D.S., Kemp, B.E. and Witters, L.A. Direct Submission Submitted (07-DEC-1995) Lee A. Witters, Medicine/Biochemistry, Dartmouth Medical School, N. College St., Hanover, NH 03755-3833, USA</p>
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JOURNAL		1. 1350
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JOURNAL MEDLINE REFERENCE		<p>2 (bases 1 to 1550) Gao, G., Widmer, J., Auster, A., Stapleton, D.S., Kemp, B.E. and Witters, L.A. Direct Submission Submitted (07-DEC-1995) Lee A. Witters, Medicine/Biochemistry, Dartmouth Medical School, N. College St., Hanover, NH 03755-3833, USA</p>
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REFERENCE		<p>1. (bases 1 to 1550) Gao, G., Fernandez, C.S., Stapleton, D., Auster, A.S., Widmer, J., Dyck, J.R., Kemp, B.E. and Witters, L.A. Non-catalytic beta- and gamma-subunit isoforms of the 5'-AMP-activated protein kinase J. Biol. Chem. 271 (15), 867</p>

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LOCUS AX281579
DEFINITION Sequence 2 from Patent WO0177305.
ACCESSION AX281579
VERSION AX281579.1 GI:16608830
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SOURCE human.
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
AUTHORS Anderson, L., Luthman, H. and Martlund, S.
TITLE Variants of the human amp-activated protein kinase gamma 3 subunit
JOURNAL Patent: WO 0177305-A 2 18-Oct-2001;
Arexis AB (SE)
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Best Local Similarity 99.3%; Pred. No. 1,3e-82;
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OY 487 gcttcgctgctgctgctgctccgcaagggcccatlcccaagctgtagagtgagagact 546
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OY 607 catggcaactgctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgct 658
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17 unordered pieces.
ACCESSION AC073128
VERSION AC073128.3 GI:13027579
KEYWORDS HTG: HTGS-PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE human.
OKANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
AUTHORS Waterston, R. H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished

REFERENCE
AUTHORS 2 (bases 1 to 196554)
TITLE Waterston, R. H.
JOURNAL Direct Submission
Submitted (08-JUN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Feb 21, 2001 this sequence version replaced g1:8469048.

COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc/index.shtml>
Project Information
Center project name: H_NH0647005
Summary Statistics
Sequencing vector: M13; 98%
Sequencing vector: plasmid; 0%
Chemistry: Dye-terminator ET; 98% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 187795 bases at least Q40
Consensus quality: 190513 bases at least Q30
Consensus quality: 192099 bases at least Q20
Insert size: 200000; agarose-fp
Insert size: 194954; sum-of-contigs
Quality coverage: 5.58 in Q20 bases; sum-of-contigs
Quality coverage: 5.67 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 17 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 1157: contig of 1157 bp in length
* 1158 1257: gap of unknown length
* 1258 3600: contig of 2343 bp in length
* 3601 3700: gap of unknown length
* 3701 5103: contig of 1403 bp in length
* 5104 5204: gap of unknown length
* 5204 8524: contig of 3321 bp in length
* 8525 8625: gap of unknown length
* 8625 11856: contig of 3232 bp in length
* 11857 11956: gap of unknown length
* 11957 15783: contig of 3827 bp in length
* 15784 15883: gap of unknown length
* 15883 21906: contig of 6023 bp in length
* 21907 22006: gap of unknown length
* 22007 28887: contig of 6881 bp in length
* 28888 28987: gap of unknown length
* 28988 35255: contig of 6268 bp in length
* 35256 35355: gap of unknown length
* 35356 44642: contig of 9287 bp in length
* 44643 44743: gap of unknown length
* 44743 58275: contig of 13533 bp in length
* 58276 58376: gap of unknown length
* 58376 73816: contig of 15441 bp in length
* 73817 73916: gap of unknown length
* 73917 92140: contig of 18224 bp in length
* 92141 92240: gap of unknown length
* 92241 113337: contig of 21097 bp in length
* 113338 113437: gap of unknown length
* 113438 130325: contig of 16888 bp in length
* 130326 130425: gap of unknown length
* 130426 149287: contig of 18662 bp in length
* 149288 149388: gap of unknown length
* 149388 196554: contig of 47167 bp in length.
Location/Qualifiers
1..196554
/organism="Homo sapiens"
/db_xref="taxon:9606"

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                  /clone="RP11-64705"
                  1..1157
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                  3701..5103
misc_feature      /note="assembly_name:Contig19
                  vector_side:right"
                  5204..8524
misc_feature      /note="assembly_name:Contig20"
                  8625..11856
misc_feature      /note="assembly_name:Contig21"
                  11957..15783
misc_feature      /note="assembly_name:Contig22"
                  15884..21906
misc_feature      /note="assembly_name:Contig23"
                  22007..28887
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                  28988..35255
misc_feature      /note="assembly_name:Contig25"
                  35356..44642
misc_feature      /note="assembly_name:Contig26"
                  44743..58275
misc_feature      /note="assembly_name:Contig27"
                  58376..73816
misc_feature      /note="assembly_name:Contig28"
                  73917..92140
misc_feature      /note="assembly_name:Contig29"
                  92241..113337
misc_feature      /note="assembly_name:Contig30"
                  113438..130325
misc_feature      /note="assembly_name:Contig31"
                  130426..149287
misc_feature      /note="assembly_name:Contig32"
                  149388..196554
misc_feature      /note="assembly_name:Contig33
                  clone_end:SP6
                  vector_side:right"
BASE COUNT      52296 a 46993 c 45889 g 49770 t 1606 others
ORIGIN
Query Match      24.7%: Score 407.2; DB 2; Length 196554;
Best Local Similarity 99.3%: Pred. No. 7.8e-83;
Matches 409; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
OY 247 agtcccccgtccagagcagctgtcgaagtcacacgggcttgagagccacatccccaagac 306
DB 62887 AGGTCCCGGTCAGCGCAGCTGTGAGTCCACCGCGCTGAGGCGCACATTCCCAAGAC 62828
OY 307 cagacccctgtcgaagctcgtccgaggttgagcctccaccaaggttgagatg 366
DB 62827 CACACCCCTGTGAAGCTGATCTGCGGGGTGGGCACTCCACCACAGGGTGGGACTG 62768
OY 367 cctcccccctgagcttaagcctcagctcaggtccagcacaagatgactgtggaagctgc 426
DB 62767 CCTCCCTCTAGCTTACAGCCTCAGCTGCAAGGCTCCAGCACAGATGATGTGGAGCTGCG 62708
OY 427 cagggagttccagcacaagagcctggagatgtgagcttaagaagcctgtctggaagaag 486
DB 62707 CACCGAGTTCCACAGCACAGAGCGCTGGAGTGTAGCTAGAAAGCCCTGTGGAAGAGAG 62648
OY 487 gcttccctgtgctgtcccccagagcccatctcccaagcttgaggttggaatgaagaaact 546
DB 62647 GCTTCCCTGTGTCTGTCTCCCGCAGGCCCAATTTCCCAAGCTGGGCTGGGATGAGAAACT 62598
OY 547 gcggaacccggcgccagatctacatgcttcatgcaagagcacaacctgtactagatgc 606
DB 62587 GCGGAACCCGCGCCCGAGATCTACATCGCTTTCAGAGGACACACTGCTTACGATGAC 62528
OY 607 catggcaactagctcagaagctagatcattctgacacacatgtcggagatcaag 658

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Db 62527 CATGGCACTAGCTCCAGGCTAGTCTTTCGACACCATCTGGAGTGAAG 62476

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RESULT 14
AF036535      1623 bp      mRNA      linear      ROD 30-MAY-2001
LOCUS      Mus musculus AMP activated protein kinase mRNA, complete cds.
DEFINITION      AF036535
ACCESSION      AF036535
VERSION      AF036535.1 GI:2766684
KEYWORDS
SOURCE
ORGANISM
house mouse.
Mus musculus.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

```

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REFERENCE
AUTHORS      1 (bases 1 to 1623)
              Shamsadin,R., Jantsan,K., Adham,I. and Engel,W.
TITLE      Cloning, organisation, chromosomal localization and expression
              analysis of the mouse Prkag1 gene
JOURNAL      Cytogenet. Cell Genet. 92 (1-2), 134-138 (2001)

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MEDLINE
PUBMED      21203559
              11306812
REFERENCE
AUTHORS      2 (bases 1 to 1623)
              Shamsadin,R.
TITLE      Direct Submission
JOURNAL      Submitted (01-DEC-1997) AG. Engel, Humangenetik, Gosslerstr.12 d,
              Goettingen 37073, Germany

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FEATURES
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Location/Qualifiers
1..1623
/organism="Mus musculus"
/db_xref="taxon:10090"
/chromosome="15"
/tissue-type="testes"
43..1035
/note="AMPK"
/codon_start=1
/product="AMP activated protein kinase"
/protein_id="AAB95475.1"
/db_xref="GI:2766685"
/translation="MESVAEESSPALLENHFOETPESSNNSVYTSFPMKSHRCYDLPTS
SKLVPDTSLOVKKAFALVTNGVRAAPRLMDSKOCFVOMLITTFDINILHRYVSAL
VQIYELERKLETWREVYLDQSKPLVCSPASSSDAYSLIRKIRLPLPYIDPESG
NTLYLTNRIRILKFLFTIEFKPEFSSLOELDGIYVANIAMVTTTPYVALGI
FVQHRVSAIPVVDENGRVVDIYSKFDVINLAAEKTVNLSVTRALXMRSHYFEGVL
KCYLHELTIIIRLVEAHVRIYVVDENHXVGLVSLDILQDLVLTGEEKRP"

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CDS

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BASE COUNT      383 a 414 c 416 g 400 t 10 others
ORIGIN
Query Match      24.1%: Score 396.8; DB 10; Length 1623;
Best Local Similarity 65.2%: Pred. No. 3e-80;
Matches 581; Conservative 0; Mismatches 310; Indels 0; Gaps 0;
OY 567 tctacatgcttcatcagagagcacacctgtctagatgcacaggaactagctcagaac 626
DB 122 TGTACTTCTCTATGATGAAGCTCATCGCTGTATGACTTAATTCACAAAGTTCCAAAGT 161
OY 627 tagcatccttcagacacatcgtgagatcaagaagcctcttctgtctgttgagcaag 686
DB 182 TGTGTGATTTGACACTTCCTGACAGGTAAAGAAAGCCTTTTGGCCCTGGTGAACAATG 241
OY 687 gctgcgggcaagccctctatggagacaagaagcagagccttctgtggagatgctgacca 746
DB 242 GTGTTCTGTGCGCGCCCTTTTGTGGACAGTAAGAGCAAGTGTGTTGTGGCATGCTGACCA 301
OY 747 tcaactgacttcatcctgtgtgtcgtacgtcactacaagtcgccctgtgtccaagatcatg 806
DB 302 TCACCGACTTATTAACAATTTTTCACCGATACCTTAAGTACAGCCCTGTGTGCAAGTTTACG 361
OY 807 agattgaacaacaataagattgagaccttgagagagatctacatcctgcaagcttccaag 866
DB 362 AACTCGAGAGCACACAGATAGAGACGTGAGAGAGAGTGTACTCTCAGAGACTCTTTAAGC 421

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 11:32:25 ; Search time 5701.1 Seconds
(without alignments)
4076.714 Million cell updates/sec

Title: US-09-826-581-3

Perfect score: 1722

Sequence: 1 cctggccctcagatcaaga.....gatgagagctcggcctgga 1722

Scoring table: IDENTITY_NUC

Gapop 10.0 ; Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues 27472414

Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estlin:*
4: em_estnu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hlc:*
9: gb_estl:*
10: gb_estc:*
11: gb_hlc:*
12: gb_gss:*
13: em_gss_hum:*
14: em_gss_inv:*
15: em_gss_pln:*
16: em_gss_vrc:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Length	ID	Description
1	148	8.6	572 10	B1344527 373008 MA
2	132.4	7.7	413 9	AA178898 zp38d10.r
3	116.8	6.8	1042 12	AL248229 Tetradion
4	82.6	4.8	933 10	BG919314 602817782
5	82.2	4.8	633 10	BC072114 BC072114
6	81.8	4.8	536 10	BM488662 pgm2n.pk0
7	80.2	4.7	595 10	BM487789 pgm2n.pk0
8	80.2	4.7	636 10	BG713637 p411n.pk0
9	80.2	4.7	647 10	BM440762 p411n.pk0
10	80.2	4.7	649 9	AJ395115
11	78.6	4.6	758 9	AJ396118
12	76.4	4.4	576 9	AV603335
13	73	4.2	564 9	AV608257
14	71	4.1	775 10	B1833269
15	70.4	4.1	448 9	AA558845
16	70.4	4.1	450 10	BP351397
17	70.4	4.1	455 9	AA578219

18	70.4	4.1	469 9	AL047390	AL047390 DKFP586A
19	70.4	4.1	473 9	BE166881	BE166881 CM4-HT050
20	70.4	4.1	508 9	BE166874	BE166874 CM4-HT050
21	70.4	4.1	583 9	AW379936	AW379936 RCA-HT025
22	70.4	4.1	586 10	BG609812	BG609812 323791 MA
23	70.4	4.1	591 9	AA410926	AA410926 FHO9C01.x
24	70.4	4.1	598 9	BE148626	BE148626 MKO-HT024
25	70.4	4.1	616 9	AW956906	AW956906 EST368976
26	70.4	4.1	634 10	BG740148	BG740148 602630747
27	70.4	4.1	668 10	BG705895	BG705895 602669396
28	70.4	4.1	708 9	AW411228	AW411228 F111B02.x
29	70.4	4.1	710 10	BI223706	BI223706 602943249
30	70.4	4.1	726 10	BI768590	BI768590 603056916
31	70.4	4.1	742 10	BI914634	BI914634 603179401
32	70.4	4.1	746 10	BF524081	BF524081 602042828
33	70.4	4.1	756 10	BI859847	BI859847 603386088
34	70.4	4.1	782 10	BI819312	BI819312 603037761
35	70.4	4.1	782 10	BI821538	BI821538 60348457
36	70.4	4.1	795 10	BE871189	BE871189 60148767
37	70.4	4.1	808 10	BI771184	BI771184 603054888
38	70.4	4.1	826 9	AL519198	AL519198 AL519198
39	70.4	4.1	841 10	BI546514	BI546514 603191524
40	70.4	4.1	864 10	BI858240	BI858240 603384001
41	70.4	4.1	870 10	BI910928	BI910928 603069326
42	70.4	4.1	903 10	BG177822	BG177822 602314222
43	70.4	4.1	905 9	AL555228	AL555228 AL555228
44	70.4	4.1	908 9	AL552459	AL552459 AL552459
45	70.4	4.1	921 9	AL548987	AL548987 AL548987

ALIGNMENTS

RESULT 1
LOCUS B1344527 572 bp mRNA linear EST 30-JUL-2001
DEFINITION 373008 MARC 2P1G Sus scrofa CDNA 5', mRNA sequence.
ACCESSION B1344527
VERSION B1344527.1 GI:15037807
KEYWORDS EST.
SOURCE pig.
ORGANISM Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
1 (bases 1 to 572)
Fahnenkrug,S.C., Freking,B.A., Rohrer,G.A., Smith,T.P.L., Casas,E.,
Stone,R.T., Heaton,M.P., Grosse,W.M., Bennett,G.A., Laegreid,W.W.
and Keele,J.W.
Design and use of two pooled tissue normalized CDNA libraries for
EST discovery in swine
Unpublished (2000)
Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt-trimmed with phred
v0.980904.e. Vector identified by cross-match with the -minscore 18
and -mismatch 12 options.

PCR Primers
FORWARD: AGGAAACGCTATGACCAT
BACKWARD: GTTTCCTCAGTCACGACG
Plate: 119 row: 1 column: 11
Seq primer: ATTTAGCTGACACTATGAG.
Location/Qualifiers

FEATURES
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1..572
/organism="Sus scrofa"
/db_xref="taxon:9823"
/clone_id="MARC 2P1G"
/csize="pooled"
/lab_host="DH10B"
/note="Vector: pCMV SPORT6; Site_1: XbaI; Site_2: XhoI;

[illegible]

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FEATURES
source
1. .933
Location/Qualifiers

RESULT 4
BC919314
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

BC919314 933 bp mRNA linear EST 05-JUN-2001
602817782p1 NC1_CGAP_Mam6 Mus musculus cDNA clone IMAGE:4946661 5'
mRNA sequence.
BC919314
BC919314.1 GI:14299790
EST.
house mouse.
Mus musculus
Eumetazoa; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 933)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: jeffrey.green@NIH.gov.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM10895 row: d column: 22
High quality sequence stop: 498.

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BASE COUNT

proving samples: century green, M.D., NIH
244 a 292 c 238 g 159 t

ORIGIN

Query Match	4.8%	Score 82.6;	DB 10;	length 933;
Best Local Similarity	52.1%;	Pred. No. 7.2e-08;		
Matches 232;	Conservative 0;	Mismatches 209;	Indels 4;	Gaps 2

Oy 1242 gccctttgaagcagcgtctcaacccttaacaaagccagatcatcgccgctgtcttg 1301
 7 gttccggattgaagctgtctcttttcatttaattggaaatTAAGATCCACAGGGCTCCAGTTATCG 66
 Oy 1302 acccggtgtcagcagaagctactcacatccctccacacacaacagcgtgtctcaagttctgc 1361
 Db 67 ACCCGAGCTCAGGCACACACCTTGACATCTCTTACTCACAAGGGATTCCTCAAGTTCTCTCA 126
 Oy 1362 acaatctttgtaag---ccgtgggcccgagtggaagaagaagggagagaccttgggcagttgac 1418
 Db 127 AGTTGTTTGAAGTAAGTAACCTTCAGGCCATCATCCGATTACCTGCGTAACATCCACAGACACAC 186
 Oy 1419 aagaagccttgagaggtcttcagcccttaagcagtcgttgaggaaagagcttggagagccctctga 1478
 Db 187 AGAGGGGTTGGGGACACAGAGGAGCGGTGGTGTGTTGATATTGAGACGTAAAGCACCTCCATCG 246
 Oy 1479 agctgcgtgagatcccttgatctccacactgtgtcccatccaaacagaggtctccctgtgtcccc 1538
 Db 247 CTCT-ATTCGGGAAAGACACCTCGCAAGCCAATGCCCCCTTCCTTCAGATWCAACGAGTTCCCA 305
 Oy 1539 ggcctctctctcccttaaccagcactatccaaagatttggagatcgtgacatctcgagaacttgg 1598
 Db 306 AGCCGGAAATTCATGTCTCTAACTCTCTCCAAAGACCTGAGATTGGCACTATGCCAATATTCG 365
 Oy 1599 ctgtgtgtgtgtgagaaagacacccatctctgacactgtgacactcttgttggaccggcgtg 1658
 Db 366 CCAATGTCCTCTACTTACACGCGCTGTCTACGTGCTGGGCACTCTTGTACAGACACCGAG 425
 Oy 1659 tgcctgcactgcctgtgtgtcaacga 1663
 Db 426 TCTCCGCTTACCTGTAAGTGAGGA 450

RESULT 5
 BJ072114
 LOCUS
 DEFINITION
 accession
 version
 keywords
 SOURCE
 ORGANISM
 XENOPUS
 laevis
 BJ072114.1
 GI:17502303
 EST.
 African clawed frog.
 Xenopus laevis
 Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Amphibia; Batrachia; Anura; Mesobatrachia; Pipridae; Pipridae;
 Xenopodinae; Xenopus.
 1 (bases 1 to 633)
 Kitayama,A., Terasaka,C., Mochii,M., Ueno,N., Shin-i,T. and Kohara
 J.
 Expressed genes in X. laevis embryo
 Unpublished (2001)
 Contact: Tadao Shin-i
 Center For Genetic Resource Information
 National Institute of Genetics
 1111 Yata, Mishima, Shizuoka 411-8540, Japan
 Tel: 81-559-81-6856
 Fax: 81-559-81-6855
 Email: tschini@genes.nig.ac.jp.
 location/Qualifiers
 1..633

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/clone_lib="NIBB mochl normalized Xenopus talbud  
library"  
/tissue_type="whole embryo"
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BASE COUNT 144 a 162 c 148 g 179 t
ORIGIN

Query Match 4.88: Score 82.2; DB 10; Length 633;
Best Local Similarity 73.4%; Pred. No. 7.4e-08;
Matches 105; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

Oy 1228 ctcccatctcgagcctgtttgaagctgtctacaccctcaagaacgagatccatcg 1287
Db 10 CTCCCGACAGACAGCCTTTTCCAGCTGTGTAATTCATCAACAATAAGATCCACCG 69
Oy 1288 cctgcctgtcttgaccggtgtcgaagcaagctacccatctccacacaaagcct 1347
Db 70 CCTACCAAGTAATGATCCCTATCTGCGACACATCTTGATCTCAGCATTAACGCGCT 129
Oy 1348 gctcaagttcctgcacatcttgaagcctcgagccca 1384
Db 130 ACTCAAGTCTCTCAACTCTTTATTGCGAGGTCCTCA 241

RESULT 6 536 bp mRNA linear EST 07-FEB-2002
LOCUS BM488662
DEFINITION pgm2n.pk008.g21 Normalized Chicken Breast Muscle, Leg Muscle, and Epiphyseal Growth Plate cDNA library (pgm2n) Gallus gallus cDNA

ACCESSION BM488662
VERSION BM488662.1 GI:18609593
KEYWORDS EST.
SOURCE chicken.
ORGANISM Gallus gallus

REFERENCE Archosauaria: Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Phasianidae; Aves; Neognathae; Galliformes; Phasianidae; Gallus. 1 (bases 1 to 536)
AUTHORS Cogburn, L.A. and Monsonego-Ornan, E.
TITLE ESTs from Normalized Chicken Breast Muscle, Leg Muscle, and Epiphyseal Growth Plate cDNA library, USDA/IFAFS Animal Genome Project

JOURNAL Unpublished (2002)
COMMENT Contact: Larry A. Cogburn
University of Delaware
Townsend Hall, Newark, DE 19717, USA
Tel: 302-831-1335
Fax: 302-831-2822
Email: cogburn@udel.edu, www.chickest.udel.edu.

FEATURES
source Location/Qualifiers
1..536
/organism="Gallus gallus"
/strain="Commercial broiler and Ottawa Res. Centre
strains 90 & 21"
/db_xref="taxon:9031"
/clone="pgm2n.pk008.g21"
/clone_1lb="Normalized Chicken Breast Muscle, Leg Muscle, and Epiphyseal Growth Plate cDNA library (pgm2n)"
/sex="Male and Female"
/tissue-type="Breast muscle, leg muscle and epiphyseal growth plate"
/dev-stage="Breast, leg: Embryo(d19): post-hatch(1d, 1, 3, 5, 7, 9, 11 weeks): growth plate(1d, 7d, 14d post-hatch)"
/lab_host="E. coli EMDH108"
/note="Vector: pCMVSPORT6; Library made from equivalent pools of total RNA isolated from each tissue (embryonic muscle 33.3%; juvenile muscle 33.3%; and epiphyseal growth plate 33.3% of the final RNA pool). Single pass sequencing from 5'-end"

BASE COUNT 117 a 171 c 132 g 116 t
ORIGIN

Query Match 4.88: Score 81.8; DB 10; Length 536;
Best Local Similarity 70.1%; Pred. No. 8.4e-08;
Matches 110; Conservative 0; Mismatches 47; Indels 0; Gaps 0;

Oy 1228 ctcccatctcgagcctgtttgaagctgtctacaccctcaagaacgagatccatcg 1287
Db 85 CTCCCGACAGCAGCCTTTTCCAGCTGTGTAATTCATCAACAATAAGATCCACCG 144
Oy 1288 cctgcctgtcttgaccggtgtcgaagcaagctacccatctccacacaaagcct 1347
Db 145 CCTCCCGTCAATCGACCCGCGACCTGCGCAACACTCTTATCATCTCAACCAACGCAAT 204
Oy 1348 gctcaagttcctgcacatcttgaagcctcgagccca 1384
Db 205 CCTCAAGTCTCTCAACTCTTTATTGCGAGGTCCTCA 241

RESULT 7 595 bp mRNA linear EST 07-FEB-2002
LOCUS BM487789
DEFINITION pgm2n.pk005.j24 Normalized Chicken Breast Muscle, Leg Muscle, and Epiphyseal Growth Plate cDNA library (pgm2n) Gallus gallus cDNA

ACCESSION BM487789
VERSION BM487789.1 GI:18608720
KEYWORDS EST.
SOURCE chicken.
ORGANISM Gallus gallus

REFERENCE Archosauaria: Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Phasianidae; Aves; Neognathae; Galliformes; Phasianidae; Gallus. 1 (bases 1 to 595)
AUTHORS Cogburn, L.A. and Monsonego-Ornan, E.
TITLE ESTs from Normalized Chicken Breast Muscle, Leg Muscle, and Epiphyseal Growth Plate cDNA library, USDA/IFAFS Animal Genome Project

JOURNAL Unpublished (2002)
COMMENT Contact: Larry A. Cogburn
University of Delaware
Townsend Hall, Newark, DE 19717, USA
Tel: 302-831-1335
Fax: 302-831-2822
Email: cogburn@udel.edu, www.chickest.udel.edu.

FEATURES
source Location/Qualifiers
1..595
/organism="Gallus gallus"
/strain="Commercial broiler and Ottawa Res. Centre
strains 90 & 21"
/db_xref="taxon:9031"
/clone="pgm2n.pk005.j24"
/clone_1lb="Normalized Chicken Breast Muscle, Leg Muscle, and Epiphyseal Growth Plate cDNA library (pgm2n)"
/sex="Male and Female"
/tissue-type="Breast muscle, leg muscle and epiphyseal growth plate"
/dev-stage="Breast, leg: Embryo(d19): post-hatch(1d, 1, 3, 5, 7, 9, 11 weeks): growth plate(1d, 7d, 14d post-hatch)"
/lab_host="E. coli EMDH108"
/note="Vector: pCMVSPORT6; Library made from equivalent pools of total RNA isolated from each tissue (embryonic muscle 33.3%; juvenile muscle 33.3%; and epiphyseal growth plate 33.3% of the final RNA pool). Single pass sequencing from 5'-end"

BASE COUNT 126 a 199 c 137 g 133 t
ORIGIN

Query Match 4.7%; Score 80.2; DB 10; Length 595;
Best Local Similarity 69.4%; Pred. No. 2e-07;
Matches 109; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 1228 ctccacatctctgagcctgtgttgaagctgtctacaccctcatcaagaacggatccatcg 1287
 DB 289 CTCCTCCCAATGCACGCTTTTGTATGCCGCTCTCCGTGATCCGCAATGAATCCACCG 348
 QY 1288 cctgcctgtctctgacccgggtgcagcaacgactccacatccctcacacaacagcct 1347
 DB 349 CTCCTCCGTATGACCGCCACCTCGGCAACACTCTCTACATCTTACACCCACAACGCAT 408
 QY 1348 gctcaagttctctgacacatcttgtaagcctgggcca 1384
 DB 409 CCTCAAGTTCTCAACTCTTTATTCAGAGTCCCA 445

RESULT 8
 LOCUS BG713637 636 bp mRNA linear EST 08-MAY-2001
 DEFINITION pgin.pk008.c13 Normalized Liver Library Gallus gallus cdna clone
 pgin.pk008.c13 5' similar to g14506061 ref|NP_002724.1| protein
 kinase, AMP-activated, gamma 1 non-catalytic subunit; AMP gamma 1;
 protein kinase, AMP-activated, noncatalytic, gamma-1 (Homo sapiens)
 g112737489 ref|XP_006778.2| protein kinase, AMP-activated, gamma 1
 non, mRNA sequence.
 ACCESSION BG713637
 VERSION HG713637.1 GI:14007587
 KEYWORDS EST.
 SOURCE Chicken.
 ORGANISM Gallus gallus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
 Phasianinae; Gallus.
 REFERENCE 1 (bases 1 to 636)
 AUTHORS Burnside, J., Morgan, R.W. and Cogburn, L.A.
 TITLE Chicken ESTs from a normalized liver library
 JOURNAL Unpublished (2001)
 COMMENT Contact: Joan Burnside
 Molecular Endocrinology
 University of Delaware
 40 Townsend Hall, Newark, DE 19717, USA
 Tel: 302 831-1345
 Fax: 302-831-3411
 Email: joan@udel.edu, www.chickest.udel.edu.
 FEATURES
 source Location/Qualifiers
 1..636
 /organism="Gallus gallus"
 /db_xref="taxon:9031"
 /clone_pgin.pk008.c13"
 /clone_lib="Normalized Liver Library"
 /sex="Male and Female"
 /tissue_type="liver"
 /lab_host="E.coli EMDH10B"
 /note="Vector: PCMVSPORT 6"
 BASE COUNT 129 a 215 c 167 g 119 t 6 others
 ORIGIN

Query Match 4.7%; Score 80.2; DB 10; Length 636;
 Best Local Similarity 69.4%; Pred. No. 2, 1e-07;
 Matches 109; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 1228 ctccacatctctgagcctgtgttgaagctgtctacaccctcatcaagaacggatccatcg 1287
 DB 429 CTCCTCCCAATGCACGCTTTTGTATGCCGCTCTCCGTGATCCGCAATGAATCCACCG 488
 QY 1288 cctgcctgtctctgacccgggtgcagcaacgactccacatccctcacacaacagcct 1347
 DB 489 CTCCTCCGTATGACCGCCACCTCGGCAACACTCTCTACATCTTACACCCACAACGCAT 548
 QY 1348 gctcaagttctctgacacatcttgtaagcctgggcca 1384
 DB 549 CCTCAAGTTCTCAACTCTTTATTCAGAGTCCCA 585

RESULT 9
 LOCUS BM440762 647 bp mRNA linear EST 01-FEB-2002
 DEFINITION pgin.pk002.19 Normalized Chicken Reproductive Tract CDNA Library
 (pgin) Gallus gallus cdna clone pgin.pk002.19 5' similar to
 g14506061 ref|NP_002724.1| protein kinase, AMP-activated, gamma 1
 non-catalytic subunit; AMP gamma 1; Protein kinase, AMP-activated,
 noncatalytic, gamma-1 (Homo sapiens) g112737489 ref|XP_006778.2|
 protein kinase, AMP-activated, gamma 1, mRNA sequence.
 ACCESSION BM440762
 VERSION BM440762.1 GI:18471537
 KEYWORDS EST.
 SOURCE Chicken.
 ORGANISM Gallus gallus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
 Phasianinae; Gallus.
 REFERENCE 1 (bases 1 to 647)
 AUTHORS Cogburn, L.A. and Nys, Y.
 TITLE ESTs from Normalized Chicken Reproductive Tract CDNA Library-
 University of Delaware and INRA, Tours-Poultry Unit Project
 JOURNAL Unpublished (2002)
 COMMENT Contact: Larry A. Cogburn
 University of Delaware
 Townsend Hall, Newark, DE 19717, USA
 Tel: 302-831-1335
 Fax: 302-831-1822
 Email: cogburn@udel.edu, www.chickest.udel.edu.
 FEATURES
 source Location/Qualifiers
 1..647
 /organism="Gallus gallus"
 /strain="Commercial broiler and layer"
 /db_xref="taxon:9031"
 /clone_pgin.pk002.19"
 /clone_lib="Normalized Chicken Reproductive Tract CDNA
 Library (pgin)"
 /sex="Male and Female"
 /tissue_type="testis, ovary and oviduct"
 /dev_stage="Various stages: embryonic, post-hatch, immature
 and sexually-mature"
 /lab_host="E. coli EMDH10B"
 /note="Vector: PCMVSPORT6; library made from three total
 RNA pools from each tissue (testis 25%, ovary 25%, and
 oviduct 50% of final RNA pool); Single pass sequencing
 from 5'-end"
 BASE COUNT 137 a 222 c 160 g 128 t

Query Match 4.7%; Score 80.2; DB 10; Length 647;
 Best Local Similarity 69.4%; Pred. No. 2, 1e-07;
 Matches 109; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 1228 ctccacatctctgagcctgtgttgaagctgtctacaccctcatcaagaacggatccatcg 1287
 DB 393 CTCCTCCCAATGCACGCTTTTGTATGCCGCTCTCCGTGATCCGCAATGAATCCACCG 452
 QY 1288 cctgcctgtctctgacccgggtgcagcaacgactccacatccctcacacaacagcct 1347
 DB 453 CTCCTCCGTATGACCGCCACCTCGGCAACACTCTTACATCTTACACCCACAACGCAT 512
 QY 1348 gctcaagttctctgacacatcttgtaagcctgggcca 1384
 DB 513 CCTCAAGTTCTCAACTCTTTATTCAGAGTCCCA 549

RESULT 10
 LOCUS AJ395115 649 bp mRNA linear EST 25-JAN-2001
 DEFINITION AJ395115 dkf426 Gallus gallus cdna 21c2rl, mRNA sequence.
 ACCESSION AJ395115
 VERSION AJ395115.1 GI:7125706
 KEYWORDS EST.

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SOURCE      chicken.
ORGANISM    gallus gallus
REFERENCE   Archosauria: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi:
AUTHORS     Archosauria: Aves: Neognathae: Galliformes; Phasianidae;
            1 (bases 1 to 649)
            Abdrakhmanov, I., Lodygin, D., Geroth, P., Atakawa, H., Law, A., Plachy
            J., Korn, B. and Buerstedde, J.M.
TITLE       A large database of chicken bursa ESTs as a resource for the
            analysis of vertebrate gene function
JOURNAL     Genome Res. 10 (12), 2062-2069 (2000)
MEDLINE    20568495
COMMENT     Contact: Buerstedde JM
            Cellular Immunology
            Heinrich-Pette-Institute
            Martinistr. 52, 20251 Hamburg, Germany
            Email: URL: http://genetics.hpi.uni-hamburg.de/dt40est.html.
FEATURES
source
    /organism="Gallus gallus"
    /strain="CB"
    /db_xref="taxon:9031"
    /clone_lib="dkfz426"
    /tissue_type="Bursa of Fabricius"

BASE COUNT      130 a      227 c      155 g      137 t
ORIGIN
Query Match      4.7%; Score 80.2; DB 9; Length 649;
Best Local Similarity 69.4%; Pred. No. 2.1e-07;
Matches 109; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

OY 1228 ctccatctctgagcctgttgaagctgtctacaccctcaagaacgagatcatcg 1287
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 46 CTCGCCCAATGCCAGCCTTTTGATGCCGTCCTCTCCGTCGCAATGAATGATCCACCG 105
OY 1288 cctgcctgtcttgacccggtgtcaggcaacgactcacatccatccacacaaacgct 1347
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 106 CTCGCCCGATCGACCGCGGCAACACTCTCTACATCTCTACCCACCAACAGCAT 165
OY 1348 gctcaagtcctgcacatcttltgaagcctggccca 1384
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 166 CCTCAAGTCTCTCAAACTCTTTATTGCGAGAGGCCCA 202

RESULT 11
LOCUS      AJ396118 758 bp mRNA linear EST 25-JAN-2001
DEFINITION AJ396118 dkfz426 Gallus gallus cDNA clone 25f16r1, mRNA sequence.
ACCESSION  AJ396118
VERSION    AJ396118.1 GI:7127728
KEYWORDS   EST.
SOURCE     chicken.
ORGANISM   Gallus gallus
            Eukaryota; Metazoa: Chordata; Craniata; Vertebrata; Euteleostomi;
            Archosauria; Aves: Neognathae; Galliformes; Phasianidae;
            Phasianinae; Gallus.
            1 (bases 1 to 758)
            Abdrakhmanov, I., Lodygin, D., Geroth, P., Atakawa, H., Law, A., Plachy
            J., Korn, B. and Buerstedde, J.M.
TITLE      A large database of chicken bursa ESTs as a resource for the
            analysis of vertebrate gene function
JOURNAL    Genome Res. 10 (12), 2062-2069 (2000)
MEDLINE    20568495
COMMENT    Contact: Buerstedde JM
            Cellular Immunology
            Heinrich-Pette-Institute
            Martinistr. 52, 20251 Hamburg, Germany
            Email: URL: http://genetics.hpi.uni-hamburg.de/dt40est.html.
FEATURES
source
    /organism="Gallus gallus"
    /strain="CB"
    /db_xref="taxon:9031"
    /clone_lib="dkfz426"
    /tissue_type="Bursa of Fabricius"

BASE COUNT      144 a      150 c      122 g      158 t
ORIGIN
Query Match      4.4%; Score 76.4; DB 9; Length 576;
Best Local Similarity 70.1%; Pred. No. 1.4e-06;

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/strain="CB"
/db_xref="taxon:9031"
/clone_lib="25f16r1"
/clone_lib="dkfz426"
/tissue_type="Bursa of Fabricius"

BASE COUNT      168 a      238 c      186 g      166 t
ORIGIN
Query Match      4.6%; Score 78.6; DB 9; Length 758;
Best Local Similarity 68.8%; Pred. No. 3e-07;
Matches 108; Conservative 0; Mismatches 49; Indels 0; Gaps 0;

OY 1228 ctccatctctgagcctgttgaagctgtctacaccctcaagaacgagatcatcg 1287
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 267 CTCGCCCAATGCCAGCCTTTTGATGCCGTCCTCTCTGATCCGCAATGAATCCACCG 326
OY 1288 cctgcctgtcttgacccggtgtcaggcaacgactcacatccatccacacaaacgct 1347
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 327 CTCGCCCGATCGACCGCGGCAACACTCTCTACATCTCTACCCCAACAGCAT 386
OY 1348 gctcaagtcctgcacatcttltgaagcctggccca 1384
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 387 CCTCAAGTCTCTCAAACTCTTTATTGCGAGAGGCCCA 423

RESULT 12
LOCUS      AV603335 576 bp mRNA linear EST 27-NOV-2001
DEFINITION AV603335 Bos taurus kidney fetus Bos taurus cDNA clone EIK1015f02
            5', mRNA sequence.
ACCESSION  AV603335
VERSION    AV603335.1 GI:9725661
KEYWORDS   EST.
SOURCE     cow.
ORGANISM   Bos taurus
            Eukaryota; Metazoa: Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidea;
            Bovidae; Bovinae; Bos.
            1 (bases 1 to 576)
            Takasuga, A., Hitotsune, S., Itoh, R., Jitchozono, A., Suzuki, H., Aso, H.
            Establishment of a high throughput EST sequencing system using
            poly(A) tail-removed cDNA libraries and determination of 36,000
            Nucleic Acids Res. 29 (22), E108 (2001)
            21570554
            Contact: Yoshikazu Sugimoto
            Animal Genetics Division
            Shirakawa Institute of Animal Genetics
            Odakura, Nishigo, Nishl-shirakawa, Fukushima 961-8061, Japan
            Tel: 81-248-25-5641
            Fax: 81-248-25-5725
            Email: kazusugie@ocn.ocn.ne.jp
            Single pass sequencing.
            This clone was obtained from a polyA-deleted cDNA library.
            Location/Qualifiers
            1..576
            /organism="Bos taurus"
            /db_xref="taxon:9913"
            /clone_lib="EIK1015f02"
            /clone_lib="Bos taurus kidney fetus"
            /tissue_type="kidney"
            /dev_stage="fetus"
            /lab_host="DH10B"
            /note="Vector: pZLI: Site_1: SalI; Site_2: NotI; Poly A
            was deleted from a NotI site"

BASE COUNT      144 a      150 c      122 g      158 t
ORIGIN
Query Match      4.4%; Score 76.4; DB 9; Length 576;
Best Local Similarity 70.1%; Pred. No. 1.4e-06;

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GAMMA CHAIN ;, mRNA sequence.
 ACCESSION AA558845
 VERSION AA388845.1 GI:2329612
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 448)
 NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)
 JOURNAL
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapds-rt@mail.nih.gov
 Tissue Procurement: W. Marston Linehan, M.D., Rodrigo Chuquai, M.D.,
 Michael R. Emmert-Buck, M.D., Ph.D.
 CDNA Library Preparation: David B. Krizman, Ph.D.
 CDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bbrp/image/image.html
 Insert length: 561 Std Error: 0.00
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 297.
 FEATURES
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 Location/Qualifiers
 1..448
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:1045913"
 /clone_11b="NCI-CGAP_Pf4.1"
 /sex="male"
 /tissue_type="prostatic intraepithelial neoplasia - high
 grade"
 /lab_host="DH10B"
 /note="Organ: prostate; Vector: PAMP10; mRNA made from
 prostatic intraepithelial neoplasia (high-grade), cDNA
 made by oligo-dT priming. Non-directionally cloned.
 Size-selected on agarose gel, average insert size 600 bp.
 CDNA Library Preparation: David B. Krizman, Ph.D.
 Reference: Krizman et al. (1996) Cancer Research
 56:5380-5383. CDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing
 Center"
 BASE COUNT 124 a 87 c 117 g 120 t
 ORIGIN
 Query Match 4.1%; Score 70.4; DB 9; Length 448;
 Best Local Similarity 68.1%; Pred. No. 2.6e-05;
 Matches 98; Conservative 0; Mismatches 46; Indels 0; Gaps 0;
 Oy 1240 cagcgtgttgaagctgtcacccatcaagaacggatcgcgtcgttct 1299
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 238 CAGCTGTGTTTATGCTGCTCTTCAATTAATTCGACACAGATCCACAGCTGCCAGTTAT 179
 Oy 1300 tgaccgggtgcaggaagctacacatccatcacacacaagcctgtcgaattcct 1359
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 178 TGACCCAGAAATACGACGATACCTGTGACATCCCTCACCCACAGGCAATCTGAAGTTCTT 119
 Oy 1360 gcaacatcttgtaagcctgggcc 1383
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 Db 118 CAATTTGTTATACACGAGTTCCC 95

Search completed: October 3, 2002, 16:14:46
 Job time: 16941 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 11:28:00 ; Search time 7316.32 Seconds

(without alignments)
4925.358 Million cell updates/sec

Title: US-09-826-581-3
Sequence: 1 cctggcccccagatcaaga.....gagagagagctcgagctgga 1722

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database:

GenEmbl: *
1: gb_ba: *
2: gb_hcg: *
3: gb_in: *
4: gb_cm: *
5: gb_ov: *
6: gb_pat: *
7: gb_ph: *
8: gb_pl: *
9: gb_pr: *
10: gb_ro: *
11: gb_sts: *
12: gb_sy: *
13: gb_un: *
14: gb_vl: *
15: em_ba: *
16: em_fun: *
17: em_hum: *
18: em_in: *
19: em_mu: *
20: em_om: *
21: em_or: *
22: em_ov: *
23: em_pat: *
24: em_ph: *
25: em_pl: *
26: em_ro: *
27: em_sts: *
28: em_un: *
29: em_vl: *
30: em_hcg_in: *
31: em_hcg_inv: *
32: em_hcg_other: *
33: em_hgq_inv: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Score	Match Length	ID	Description
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1	1722	100.0	1722	6	AX281580	Sequence
2	1719.4	99.8	152129	2	AC027416	Homo sapi
3	1697	98.5	196554	2	AC073128	Homo sapi
4	1697	98.5	206854	4	AC009974	Homo sapi
5	612.6	35.6	5888	4	AF214521	Homo sapi
6	209.4	12.2	227224	2	AF336381	Sus scrofa
7	168.2	9.8	1647	6	AX281582	Mus muscu
8	168.2	9.8	2109	6	AX099776	Sequence
9	168.2	9.8	2115	6	AX099802	Sequence
10	168.2	9.8	2115	6	AF214519	Sequence
11	168.2	9.8	2290	6	HS249977	Sequence
12	148	8.6	1867	6	AX099774	Homo sapi
13	148	8.6	1873	4	AX214520	Sequence
14	148	8.6	1873	6	AX099800	Sequence
15	148	8.6	2022	6	AX099804	Sequence
16	91.4	5.3	106	11	G67375	Sequence
17	82.2	4.8	142903	2	AL627254	Sequence
18	79.4	4.6	14411	4	AF329081	Sequence
19	77	4.5	73638	2	AC015613	Sequence
20	74.2	4.3	81210	2	AC019242	Sequence
21	74.2	4.3	171822	2	AC011603	Sequence
22	74.2	4.3	180749	2	AC025256	Sequence
23	70.4	4.1	1578	9	HS042412	Sequence
24	70.4	4.1	1677	9	BC000358	Sequence
25	69.4	4.0	101215	9	AC006966	Sequence
26	68	3.9	3497	3	AF094763	Sequence
27	68	3.9	80069	2	AC019671	Sequence
28	68	3.9	173634	3	AC009344	Sequence
29	68	3.9	195868	3	AC008308	Sequence
30	68	3.9	230266	3	AE003733	Sequence
31	67.6	3.9	7218	6	166494	Sequence
32	65.8	3.8	2082	3	AF094764	Sequence
33	65.8	3.8	4666	3	AY070541	Sequence
34	65.6	3.8	104	10	AF266988	Sequence
35	65.6	3.8	1328	10	RNAMPKGM	Sequence
36	65.6	3.8	1550	10	RNMD2413	Sequence
37	65.6	3.8	1623	10	AF036535	Sequence
38	63.8	3.7	1167	9	AB025580	Sequence
39	63.8	3.7	1435	6	AR139104	Sequence
40	63.8	3.7	2062	9	HS249976	Sequence
41	63.8	3.7	2194	9	AF087875	Sequence
42	63.8	3.7	2203	9	BC020540	Sequence
43	63.8	3.7	2223	9	AK001887	Sequence
44	63.8	3.7	3132	10	BC015283	Sequence
45	62.4	3.6	125020	9	AF429315	Sequence

ALIGNMENTS

RESULT	1
AX281580	Sequence 3 from Patent WO0177305.
LOCUS	AX281580
DEFINITION	Sequence 3 from Patent WO0177305.
ACCESSION	AX281580
VERSION	AX281580.1 GI:16608831
KEYWORDS	
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
TITLE	Andersson, L., Luthman, H. and Marklund, S.
JOURNAL	Patent: WO 0177305-A 3 18-Oct-2001;
AREXIS	Arexis AB (SE)
FEATURES	Location/Qualifiers
source	1..1722
BASE COUNT	321 a 504 c 534 g 363 t
ORIGIN	/organism="Homo sapiens"
	/db_xref="taxon:9606"

TITLE
JOURNAL
COMMENT

Grand-Pierre, N., Granl, G., Hagos, B., Heaford, A., Horton, L.,
Kiehl, J. C., Iliev, I., Johnson, R., Jones, C., Kann, L., Karatas, A.,
Klein, J., Lakoque, K., Lamazares, R., Landers, T., Lehoczy, J.,
Levine, R., Lieu, C., Liu, C., Locke, K., Macdonald, P., Margolis, N.,
McCarthy, M., McKean, P., McCurk, A., McKernan, K., McPheters, R.,
Meldrum, J., Menus, L., Mihova, T., Miranda, C., Mlenga, V., Morrow, J.,
Murphy, T., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P.,
O'Neill, D., Oliver, T. M., Oliver, J., Peterson, K., Pierre, N.,
Pisani, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D.,
Roy, A., Santos, R., Schauer, S., Sever, P., Spencer, B.,
Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,
Testaye, S., Theodore, J., Tirrell, A., Travers, M., Triggillo, J.,
Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J.,
Young, G., Zainoun, J., Zimmer, A. and Zody, M.

Direct Submission
Submitted (30-MAR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 7, 2000 this sequence version replaced gi:7342115.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: MIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu

Project Information
Center project name: L7458
Center clone name: 504_G_11

Summary Statistics
Sequencing vector: M13: M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 135376 bases at least Q40
Consensus quality: 143264 bases at least Q30
Consensus quality: 146503 bases at least Q20
Insert size: 161000; agarose-fp
Insert size: 149029; sum-of-ctrls
Quality coverage: 3.1 in Q20 bases; agarose-fp
Quality coverage: 3.3 in Q20 bases; sum-of-ctrls

NOTE: This is a 'working draft' sequence. It currently
consists of 32 contigs. The true order of the pieces
is not known and their order in this sequence record is
arbitrary. Gaps between the contigs are represented as
runs of N, but the exact sizes of the gaps are unknown.
This record will be updated with the finished sequence
as soon as it is available and the accession number will
be preserved.

1. 1005: contig of 1005 bp in length
1006 1105: gap of 100 bp
1106 2402: contig of 1297 bp in length
2403 2502: gap of 100 bp
2503 3823: contig of 1321 bp in length
3824 3923: gap of 100 bp
3924 5020: contig of 1097 bp in length
5021 5120: gap of 100 bp
5121 6161: contig of 1041 bp in length
6162 6261: gap of 100 bp
6262 7547: contig of 1286 bp in length
7548 7647: gap of 100 bp
7648 9883: contig of 2336 bp in length
9884 10083: gap of 100 bp
10084 12556: contig of 2473 bp in length
12557 12656: gap of 100 bp
12657 15043: contig of 2387 bp in length
15044 15143: gap of 100 bp
15144 17123: contig of 1960 bp in length
17124 17223: gap of 100 bp
17224 19466: contig of 2243 bp in length
19467 19566: gap of 100 bp
19567 21928: contig of 2362 bp in length
21929 22028: gap of 100 bp

FEATURES

22029 24319: contig of 2291 bp in length
24320 24419: gap of 100 bp
24420 27059: contig of 2640 bp in length
27060 27159: gap of 100 bp
27160 30170: contig of 3011 bp in length
30171 30270: gap of 100 bp
30271 33968: contig of 3698 bp in length
33969 34068: gap of 100 bp
34069 38179: contig of 4111 bp in length
38180 38279: gap of 100 bp
38280 42366: contig of 4087 bp in length
42367 42466: gap of 100 bp
42467 46365: contig of 3899 bp in length
46366 46465: gap of 100 bp
46466 51285: contig of 4820 bp in length
51286 51385: gap of 100 bp
51386 55871: contig of 4486 bp in length
55872 55971: gap of 100 bp
55972 60595: contig of 4624 bp in length
60596 60695: gap of 100 bp
60696 66595: contig of 5900 bp in length
66596 66695: gap of 100 bp
66696 73218: contig of 6523 bp in length
73219 73318: gap of 100 bp
73319 77115: contig of 3797 bp in length
77116 77215: gap of 100 bp
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85023 85122: gap of 100 bp
85123 93314: contig of 8192 bp in length
93315 93414: gap of 100 bp
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101194 101293: gap of 100 bp
101294 113090: contig of 11797 bp in length
113091 113190: gap of 100 bp
113191 123496: contig of 10306 bp in length
123497 123596: gap of 100 bp
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QY 121	ggagatgtagagaggtgtgagggggagatctgtacggttgtctcgtggctgatactgatat	180		
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QY 181	accacaaagcttggtctcgaagccaagcccacaggggccaaggtgtgagaaagctcatcc	240		
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QY 241	ggagctcgtgcatlgtgcagagctcggagagaccctcggggtctcaaltctcccatctgtgagccgt	300		
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QY 301	atgaccagctgacaccccttccactccgtactgcatlgtgacctgtgccaagtgtctaa	360		
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QY	1201	ctccctaggcgtgcccgaggtcgaactgtctccacatctctgaagcctgtttgaagctgtcta	1260
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QY	1441	ccctgaagctgttggggaagagcttggaagccctctgaagctgtcgtgatactctgaatccc	1500
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OY 1681 CCAATGTCGTACCCACCCAGATGAGAGGCTCGGCTGG 1721
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Locus Homo sapiens chromosome 2 clone RP11-64705, WORKING DRAFT SEQUENCE,
DEFINITION 17 unordered pieces.
AC073128 AC073128.3 GI:13027579
VERSION HTG: HTGS_PHASE1: HTGS_DRAFT: HTGS_FULLTOP.
KEYWORDS human.
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 196554)
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
AUTHORS 2 (bases 1 to 196554)
TITLE Waterston,R.H.
JOURNAL Direct Submission
AUTHORS Submitted (08-JUN-2000) Genome Sequencing Center, Washington
TITLE University School of Medicine, 4444 Forest Park Parkway, St. Louis,
JOURNAL MO 63108, USA
COMMENT On Feb 21, 2001 this sequence version replaced gi:8469048.
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
Project Information -----
Center project name: H_NH0647005
Summary Statistics -----
Sequencing vector: MJ3; 988
Sequencing vector: plasmid; 08
Chemistry: Dye-terminator Big Dye; 08 of reads
Chemistry: Dye-terminator Big Dye; 08 of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 18705 bases at least Q40
Consensus quality: 190513 bases at least Q30
Consensus quality: 192099 bases at least Q20
Insert size: 200000; agarose-fp
Insert size: 194954; sum-of-contigs
Quality coverage: 5.58 in Q20 bases; agarose-fp
Quality coverage: 5.67 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 17 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 1157: contig of 1157 bp in length
* 1158 1257: gap of unknown length
* 1258 3600: contig of 2343 bp in length
* 3601 3700: gap of unknown length
* 3701 5103: contig of 1403 bp in length
* 5104 5203: gap of unknown length
* 5204 8524: contig of 3321 bp in length
* 8525 8624: gap of unknown length
* 8625 11856: contig of 3232 bp in length
* 11857 11956: gap of unknown length
* 11957 15783: contig of 3827 bp in length
* 15784 15883: gap of unknown length
* 15884 21906: contig of 6023 bp in length

* 21907 22006: gap of unknown length
* 22007 28887: contig of 6881 bp in length
* 28888 28987: gap of unknown length
* 28988 35255: contig of 6268 bp in length
* 35256 35356: gap of unknown length
* 35356 44642: contig of 9287 bp in length
* 44643 44742: gap of unknown length
* 44743 58275: contig of 13533 bp in length
* 58276 58375: gap of unknown length
* 58376 73817: contig of 15441 bp in length
* 73817 73916: gap of unknown length
* 73917 92140: contig of 18224 bp in length
* 92141 92240: gap of unknown length
* 92241 113337: contig of 21097 bp in length
* 113338 113437: gap of unknown length
* 113438 130325: contig of 16888 bp in length
* 130326 130425: gap of unknown length
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* 149288 149387: gap of unknown length
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Best Local Similarity 99.9% Pred. No. 0;
Matches 1719; Conservative 0; Mismatches 0; Indels 2; Gaps 2;
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[illegible]

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Oy	1621	catcctgactgactcgcggacatcttgtgagccggcggtgtgtctgcaactgcctgtgtcaa	1680
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DEFINITION	Homo sapiens BAC clone RP11-459119 from 2, complete sequence.		PRI 09-JAN-2002
ACCESSION	AC009974		
VERSION	AC009974.9	GI:16799058	
KEYWORDS	HTG.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
TITLE	1 (bases 1 to 206854)		
JOURNAL	Sullivan,J.E. and Waterston,R.		
MEDLINE	Toward a complete human genome sequence		
REFERENCE	Genome Res. 8 (11), 1097-1108 (1998)		
AUTHORS	2 (bases 1 to 206854)		
TITLE	Harris,A. and Cotton,M.		
JOURNAL	The sequence of Homo sapiens BAC clone RP11-459119		
REFERENCE	unpublished (2001)		
AUTHORS	3 (bases 1 to 206854)		
TITLE	Waterston,R.H.		
JOURNAL	Direct Submission		
REFERENCE	Submitted (08-SEP-1999) Genome Sequencing Center, Washington		
AUTHORS	University School of Medicine, 4444 Forest Park Parkway, St. Louis,		
TITLE	MO 63108, USA		
JOURNAL	4 (bases 1 to 206854)		
REFERENCE	Waterston,R.H.		
AUTHORS	Direct Submission		
TITLE	Submitted (08-NOV-2001) Genome Sequencing Center, Washington		
JOURNAL	Submitted (08-NOV-2001) Genome Sequencing Center, Washington		

University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE
5 (bases 1 to 206854)
AUTHORS
Waterston,R.H.
TITLE
Direct Submission
JOURNAL
Submitted (03-JAN-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE
6 (bases 1 to 206854)
AUTHORS
Waterston,R.
TITLE
Direct Submission
JOURNAL
Submitted (09-JAN-2002) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Nov 8, 2001 this sequence version replaced gi:13431203.

COMMENT
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc>
Contact: saplens@watson.wustl.edu
----- Summary Statistics
Center project name: H_NH0459119

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Woon,P.Y., Zhao,B., Frengen,E., Tateo,M., Calanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Koswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6
NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is RP11-1077K22: the clone sequenced to the right is RP11-64705. Actual start of this clone is at base position 1 of RP11-459119; actual end is at base position 206854 of RP11-459119.

Data from AC079810 and AC073128 was used to finish this clone, AC009974. Polymorphisms have been identified between AC073128 and AC009974. A single plasmid region exists between 38812-38903. An unresolved tandem in the HERV SVA exists between 184390-185163. PCR suggests that approximately 1700 bps are missing.

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			Gaps	2;

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QY	1441	cccttagcaatctgggggaagaagcttggaagccctctgaaagctctgtgtgatactctctc	1500
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QY	1501	acctgtgccccaactcaaaccaaggtctccctgctgcctccggcctcctcctctcaacgac	1560
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QY	1561	tatccaagaatttgggcatcggcgaattctccggagactttggctgtgtgtctgtagaagaacacc	1620
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QY	1681	cgaatgtgtgtacccaacccccaagatgaagaagctctggagctgg	1721
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RESULT 5 AF214521 5888 bp DNA linear MAM 03-JUN-2000
LOCUS Sus scrofa AMPK gamma subunit (PRKAG3) gene, complete cds.
DEFINITION AF214521
ACCESSION AF214521 GI:8215685
VERSION
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suidae; Sus.
REFERENCE 1 (bases 1 to 5888)
AUTHORS Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Thelander,M.,
Kogel-Galliard,C., Paul,S., Gellin,J., Lundstrom,K., Reinsch,N.,
Kogel-Galliard,C., Paul,S., Iannucelli,N., Rask,L., Rome,H.,
Lundstrom,K., Reinsch,N., Gellin,J., Kalm,E., Roy,P.L., Chardon,P.
and Andersson,L.
A mutation in PRKAG3 associated with excess glycogen content in pig
skeletal muscle
Science 288 (5469), 1248-1251 (2000)
JOURNAL MEDLINE 20280150
PUBMED 10818001
REFERENCE 2 (bases 1 to 5888)
AUTHORS Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A.,
Kogel-Galliard,C., Paul,S., Gellin,J., Lundstrom,K., Reinsch,N.,
Kalm,E., Le Roy,P., Chardon,P. and Andersson,L.
Direct Submission
Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish
University of Agricultural Sciences, BMC Box 597, Uppsala 751 24,
Sweden
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Best Local Similarity 66.3%; Pred. No. 1e-137;
Matches 1155; Conservative 0; Mismatches 484; Indels 102; Gaps 15;

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Oy	1292	ccgtctcttgaccggcgtgtcagaacgtaacctcacatctccacacacaaagcgtctc	1351
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Oy	1470	ccctcttgaaagctgtgataccctgatctcaacctgtgtcccatctaaccaaggctccc	1529
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Oy	1530	lactgccccggagccctccttctctctacagacatccaagaattggacalcgacatccc	1589
Dn	3293	TCTGTCCCCGGCCCTCTCTTCTACCCGACCATCAAMATTTGGCATGGCAATTC	3352
Oy	1590	gaagacttggcttgggtgcttgagaaagacaccatctgaacttgaagcatcttctg	1649
Dn	3353	GAGACTGTGGCCCTGGTGCTGAAGACGGGCCCATCTGTGACCGCACTGGACATCTTCGTGG	3412
Oy	1650	accggcggtgtgctgcatctgctgtgtgtcaagaatgtgtgtacccaccagatgaga	1709
Dn	3413	ACCGGCTGTGTCTGCCCTGTGTCAACGAACGTAACTATGCCAGATGGGG	3472
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Dn	3473	G 3473	
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DEFINITION	Mus musculus chromosome 1 clone PAC510; PAC457, *** SEQUENCING IN PROGRESS ***, 3 unordered pieces.		
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VERSION	AF336381.1	GI:13507298	
KEYWORDS	HTG; HTGS-PHASE1.		
SOURCE	house mouse.		
ORGANISM	Mus musculus.		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.		
AUTHORS	Rump,A., Hayes,C., Brown,S.D.M. and Rosenthal,A.		
JOURNAL	Mouse chromosome 1 genomic sequence unpublished		
REFERENCE	2 (bases 1 to 227724)		
TITLE	Rump,A.		
AUTHORS	Direct Submission		
JOURNAL	Submitted (17-JAN-2001) Genome Analysis, Institute of Molecular Biotechnology, Beutenbergstr. 11, Jena 07745, Germany		
COMMENT	* NOTE: This is a 'working draft' sequence. It currently consists of 3 contigs. The true order of the pieces * is not known and their order in this sequence record is * arbitrary. Gaps between the contigs are represented as * runs of N, but the exact sizes of the gaps are unknown. * This record will be updated with the finished sequence * as soon as it is available and the accession number will * be preserved.		

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TITLE	Variants of the human amp-activated protein kinase gamma 3 subunit				
JOURNAL	Patient: WO 0177305-A 5 18-OCT-2001;				
	Arexis AB (SE)				
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ORIGIN					
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Best Local Similarity	95.6%; Pred. No. 4,2e-30;				
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Oy	1570	tttgggcatcggaacatccgagacttgacgtggtgctgtagagacagaccatcctgac	1629		
Db	1069	TTTGGGCATCCGGCACATCCAGACATTGGCTGTGGTGTGGAGACAGCACCATCTGAC	1128		
Oy	1630	tgcactgacacatcttttgagacggcgagtgctgacactgacctgtgtcaacgaatctgg	1689		
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DEFINITION	Sequence 3 from Patent WO0120003.				
ACCESSION	AX099776				
VERSION	AX099776.1 GI:13538810				
KEYWORDS					
SOURCE	human.				
ORGANISM	Homo sapiens				
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.				
REFERENCE	1 (bases 1 to 2109)				
AUTHORS	Andersson,L., Looft,C., Kalm,E., Milan,D., Robic,A., Rogel-Galliard,C., Iannuccielli,N., Gellin,J., Le Roy,P. and Charodon,P.				
TITLE	Variants of the gamma chain of ampk, dna sequences encoding the same, and uses thereof				
JOURNAL	Patent: WO 0120003-A 3 22-MAR-2001; INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ; Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)				
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Matches 160: Conservative 0; Mismatches 20; Indels 0; Gaps 0;

QY 1510 ccacttaacaggggttcctctgctgcccgcgcctctctctctctacgcacatccaga 1569
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QY 1570 ttgggcatcggcacatccgagacttgctgctgctgagagacacacatctgac 1629
DB 975 TTTGGGCAATCGGCACATTCGAGACTTGCGCTGTGGAAGGCGCCCATCTGCAC 1034
QY 1630 tgcactgacatcttcttgaccggcgctgctgacactccctgctgtaacgaatgtg 1689
DB 1035 CGCACTGACATCTTCTGACCGCGCTGTGTGCGCTGCTGCTGTGCTGAACGAACCTGG 1094

RESULT 14
LOCUS AX099800 1873 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 27 from Patent WO0120003.
ACCESSION AX099800
VERSION AX099800.1 GI:13538834
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Sus.
1 (bases 1 to 1873)
Rogel-Galliard, C., Iannuccelli, N., Gellin, J., Le Roy, P. and
Charbon, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 27 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kaim, Ernst (DE)
Location/Qualifiers
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/db_xref="taxon:9823"
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AVULETAPILALDIFVDRVSALPVNVEGTVGLYSRFVTHLAQOQTYNHLDMY
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BASE COUNT 382 a 580 c 535 g 376 t
ORIGIN

Query Match 8.6%; Score 148; DB 6; Length 1873;
Best Local Similarity 88.9%; Pred. No. 3.2e-25;
Matches 160: Conservative 0; Mismatches 20; Indels 0; Gaps 0;

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QY 1570 ttgggcatcggcacatccgagacttgctgctgctgagagacacacatctgac 1629
DB 975 TTTGGGCAATCGGCACATTCGAGACTTGCGCTGTGGAAGGCGCCCATCTGCAC 1034
QY 1630 tgcactgacatcttcttgaccggcgctgctgacactccctgctgtaacgaatgtg 1689
DB 1035 CGCACTGACATCTTCTGACCGCGCTGTGTGCGCTGCTGCTGTGCTGAACGAACCTGG 1094

RESULT 15
LOCUS AX099804 2022 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 31 from Patent WO0120003.
ACCESSION AX099804
VERSION AX099804.1 GI:13538838
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Sus.
1 (bases 1 to 2022)
Rogel-Galliard, C., Iannuccelli, N., Gellin, J., Le Roy, P. and
Charbon, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 31 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kaim, Ernst (DE)
Location/Qualifiers
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/organism="Sus scrofa"
/db_xref="taxon:9823"
593 g 394 t

BASE COUNT 412 a 623 c 593 g 394 t
ORIGIN

Query Match 8.6%; Score 148; DB 6; Length 2022;
Best Local Similarity 88.9%; Pred. No. 3.1e-25;
Matches 160: Conservative 0; Mismatches 20; Indels 0; Gaps 0;

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QY 1570 ttgggcatcggcacatccgagacttgctgctgctgagagacacacatctgac 1629
DB 1125 TTTGGGCAATCGGCACATTCGAGACTTGCGCTGTGGAAGGCGCCCATCTGCAC 1184
QY 1630 tgcactgacatcttcttgaccggcgctgctgacactccctgctgtaacgaatgtg 1689
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Search completed: October 3, 2002, 14:46:08
Job time: 11888 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 12:32:10 ; Search time 719.93 Seconds
(without alignments)
4106.686 Million cell updates/sec

Title: us-09-826-581-3

Perfect score: 1722

Sequence: 1 cctggccctcagatcaaga.....gatgagagctcggtcgtga 1722

Scoring table:

IDENTITY_NUC
Gapox 10.0, Gapexl 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1722	100.0	1722	22	AAH43683
2	351	20.4	378	22	ABAA44706
3	351	20.4	378	22	ABA55162
4	351	20.4	378	22	ABA24907
5	351	20.4	378	22	AAK03423
6	351	20.4	378	22	AAK28874
7	351	20.4	378	22	AAI13465
8	351	20.4	378	22	AAI34821
9	351	20.4	378	22	AAI03344

10	196	11.4	547	22	ABA08485	Human AMP-activate
11	168.2	9.8	1647	22	AAH43685	PRKAG3 CDNA. Homo
12	168.2	9.8	2109	22	AAD03296	Human AMPK gamma s
13	168.2	9.8	2115	22	AAD03320	Human AMPK gamma s
14	148	8.6	1867	22	AAD03295	Pig AMPK gamma sub
15	148	8.6	1873	22	AAD03319	Pig AMPK gamma sub
16	148	8.6	2022	22	AAD03321	Sus scrofa PRKAG3
17	74.2	4.3	16525	22	AAK73303	Human immune/haema
18	70.4	4.1	602	22	AAH35203	Human colon cancer
19	70.4	4.1	1691	21	AAC98774	Human pancreatic c
20	68.8	4.0	1576	18	AAT85927	Mammalian AMPK-gam
21	68	3.9	39651	23	ABL18856	Drosophila melanog
22	65.8	3.8	3261	23	ABL18857	Drosophila melanog
23	65	3.8	92	22	ABA49850	Human breast cell
24	65	3.8	92	22	ABA67769	Human foetal liver
25	65	3.8	92	22	ABA4826	Probe #13292 for g
26	65	3.8	92	22	AAK16181	Human brain expres
27	65	3.8	92	22	AAK41922	Human bone marrow
28	65	3.8	92	22	AAI22692	Probe #12625 for g
29	65	3.8	92	22	AAI47988	Probe #16674 used t
30	65	3.8	92	22	AAI08354	Probe #345 used t
31	63.8	3.7	1435	20	AAK06882	Disease associated
32	63.8	3.7	1467	23	AAK84265	DNA encoding novel
33	63.8	3.7	2223	22	AAH14839	Human CDNA sequenc
34	62.6	3.6	350	21	AAC01661	Human secreted pro
35	57.2	3.3	735	22	AAH07561	Human CDNA clone (
36	49.8	2.9	2303	23	AAK84267	DNA encoding novel
37	49.4	2.9	151	24	AB199496	Mouse ischaemic co
38	44	2.6	375	22	AAK6844	Novel human polynu
39	42.6	2.5	985	22	AAK08652	Mouse cancer assoc
40	42	2.4	61710	22	AAK83782	Human immune/haema
41	41.6	2.4	37664	22	AAK83781	Human immune/haema
42	40.8	2.4	1512	21	AAK288024	Human secreted pro
43	40.8	2.4	1512	21	AAK1637	Human secreted pro
44	40.6	2.4	766	21	AAK4215	Arabidopsis thalia
45	40.6	2.4	8017	22	ABAI9015	Human nervous syst

ALIGNMENTS

RESULT 1	AAH43683	standard; DNA; 1722 BP.
ID	AAH43683	
XX	AAH43683;	
AC	21-JAN-2002	(first entry)
XX	PRKAG3 intron 4 - intron 10.	
DE	Human; AMP-activated protein kinase gamma 3 subunit; PRKAG3; variant;	
KW	metabolic disease; diabetes; obesity; substitution; ds.	
KM		
XX	Homo sapiens.	
OS		
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FT	exon	14..95
FT	exon	/*tag- b
FT	intron	/number- "Exon 5"
FT	intron	96..552
FT	intron	/*tag- c
FT	exon	/number- "intron 5"
FT	exon	553..611
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FT      /number= "Intron 10"
FT      /note= "5' portion of Intron 10"
PN      WO200177305-A2.
XX      18-OCT-2001.
PD      06-APR-2001; 2001WO-SE00765.
XX      07-APR-2000; 2000US-195665P.
PR      (AREX-) AREXIS AB.
XX      Andersson L, Luthman H, Marklund S;
DR      WPI: 2001-657170/75.
XX      New variants of human AMP-activated protein kinase gamma3 subunit
PT      associated with a metabolic disease e.g. diabetes or obesity and method
PT      for determining a risk estimate of diseases in subject by detecting the
PT      variant -.
XX      Example 1; Fig 3; 25pp; English.
PS      The sequences given in AAH43681-84 represents genomic fragments
CC      encoding the human AMP-activated protein kinase gamma 3 subunit
CC      (PRKAG3). Detecting the presence of the PRKAG3 DNA, or a variant,
CC      is useful in determining a risk estimate of a metabolic disease,
CC      such as diabetes or obesity, in a subject. The variation may occur
CC      in exons 3, 4 or 10. In exon 3 variation may be a substitution of
CC      a G for a C at nucleotide 320, resulting in the amino acid
CC      substitution P71A: in exon 4 variation may be a substitution of a
CC      T for a C at nucleotide 550, and in exon 10 variation may be a
CC      substitution of a T for a C at nucleotide 1037, resulting in the
CC      amino acid substitution R340W. There may also be nucleotide variation
CC      in Intron 6.
XX      Sequence 1722 BP: 321 A: 504 C: 534 G: 363 T: 0 other;
SO      Query Match 100.0%; Score 1722; DB 22; Length 1722;
      Best Local Similarity 100.0%; Pred. No. 0;
      Matches 1722; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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AC			
XX			
DT	01-FEB-2002	(first entry)	
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DE	Human breast cell single exon nucleic acid probe #3401.		
KW	Human; microarray: single exon probe; gene expression: breast;		
KW	disease: cancer; ss.		
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OS	Homo sapiens.		
XX			
PN	W0200157271-A2.		
PD			
XX	09-AUG-2001.		
PF	30-JAN-2001; 2001MO-US00662.		
XX			
PR	04-FEB-2000; 2000US-0180312.		
PR	26-MAY-2000; 2000US-0207456.		
PR	30-JUN-2000; 2000US-0608408.		
PR	03-AUG-2000; 2000US-0633366.		
PR	21-SEP-2000; 2000US-0234687.		
PR	27-SEP-2000; 2000US-0236359.		
PR	04-OCT-2000; 2000GB-0024263.		
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PA	(MOLE-) MOLECULAR DYNAMICS INC.		
XX			
PI	Penn SG, Hanzel DK, Chen W, Rank DR;		
XX			
PR	WPI; 2001-496933/54.		
XX			

PT New spatially-addressable set of single exon nucleic acid probes,
PT useful for measuring gene expression in sample derived from
PT breast, comprises number of single exon nucleic acid probes -
XX
XX
PS Claim 1; SEQ ID NO 3401; 327bp + sequence listing; English.
XX
CC The invention relates to a spatially-addressable set of single exon
CC nucleic acid probes for measuring gene expression in a sample derived
CC from human breast and BT 474 cells. The method involves contacting
CC the probes with a collection of detectably labelled nucleic acids
CC derived from mRNA of human breast, and then measuring the label
CC bound to each probe of the microarray. The probes are useful for
CC verifying the expression of regions of genomic DNA predicted to
CC encode proteins. They are useful for gene discovery, and for
CC determining predisposition and/or prognosing breast disease. Gene
CC expression analysis is useful for assessing the toxicity of chemical
CC agents on cells. The microarray of this invention presents a far greater
CC diversity of probes for measuring gene expression, with far less bias
CC than expressed sequence tag microarrays. The method is suitable for
CC rapid production of functional information from genomic sequence. The
CC present sequence is a single exon nucleic acid probe of the invention.
CC Note: The sequence data for this patent did not form part of the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at fcp.wipo.int/pub/published_pat_sequences.

SQ Sequence 378 BP; 80 A; 128 C; 97 G; 73 T; 0 other;

Query Match	20.48; Score 351; DB 22; Length 378;
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Best Local Similarity 99.7%; Pred. NO. 6.9e-83;
Matches 362; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

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db 242 GGAGATGGAGGAGGTGAGGGGAGATCTTTGTACGGTCTTCTGGGGCTGATCTGATAT 183

OV 181 accacaaqcttgcattcaqqccaaqccaqqcgaqqqtgaaaatccatcc 240

Db 182 ACCACAAGCTTGGCTTCAGGCCAAGCCAGGCCAGGGCCAGGCTGAGGAAAGTCATCC 123

OY 241 gga gtc tgc atg gcc agc tgg gga gac ccctgg ggc tca attcc catctgttg agccgct 300

Db 122 GGAGTCTGCATGGCCAGCTGGGAGACCCCTGGGCTCAATTCCCCATCTGTGGAGCCGCT 63

301 atgacca gctgacac cttca cctccg cta ctgcat ggcctg tgcata ggtgtc tagg 360

Db 62 ATGACCAGCTGACACCTTTCACCTCCGCTACTGCATGGCCCTGTC-CATAGGTCCTAGGG 4

Qy 361 agc 363
111

Db 3 AGC 1

RESULT 3

ABA55162/C
ID ABA55162 standard: DNA: 378 BP.

XX
AC ABA55162;

AA	
DT	01-FEB-2002 (first entry)

AA Human foetal liver single exon nucleic acid probe #3467.

Human; foetal liver; gene expression; single exon nucleic acid probe; ss

OS Homo sapiens.
XX ABA24907/C
XX ID ABA24907 standard: DNA: 378 BP.
XX
XX
XX PD 09-AUG-2001.
XX
XX PF 30-JAN-2001: 2001WO-US00669.
XX
XX PR 04-FEB-2000: 2000US-0180312.
XX PR 26-MAY-2000: 2000US-0207456.
XX PR 30-JUN-2000: 2000US-0608408.
XX PR 03-AUG-2000: 2000US-0632366.
XX PR 21-SEP-2000: 2000US-0234687.
XX PR 27-SEP-2000: 2000US-0236359.
XX PR 04-OCT-2000: 2000GB-0024263.
XX
XX PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX PI Penn SG, Hanzel DK, Chen W, Rank DR:
XX
XX DR WPI: 2001-483447/52.
XX
XX PT Human genome-derived single exon nucleic acid probes useful for
XX PT analyzing gene expression in human fetal liver -
XX
XX PS Claim 1: SEQ ID NO 3467: 639pp + sequence listing: English.
XX
XX CC The invention relates to a single exon nucleic acid probe for
XX CC measuring human gene expression in a sample derived from human foetal
XX CC liver. The single exon nucleic acid probes may be used for predicting,
XX CC measuring and displaying gene expression in samples derived from human
XX CC fetal liver. The present sequence is a single exon nucleic acid
XX CC probe of the invention.
XX CC Note: The sequence data for this patent did not form part of the
XX CC printed specification, but was obtained in electronic format directly
XX CC from WIPO at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX SQ Sequence 378 BP; 80 A; 128 C; 97 G; 73 T; 0 other:

Query Match 20.4%; Score 351; DB 22; Length 378;
Best Local Similarity 99.7%; Pred. No. 6.9e-83;
Matches 362; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
DB 1 cctggccctcagatcaagaagcctcttctgtctgtgagcgaaggtgtgagcagc 60
DB 362 cctggccctcagatcaagaagcctcttctgtctgtgagcgaaggtgtgagcagc 303
OY 61 cctctatgagcagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 120
DB 302 cctctatgagcagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 243
OY 121 ggaagatgagcgaagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 180
DB 242 ggaagatgagcgaagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 183
OY 181 accacaagccttgcctcagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagc 240
DB 182 accacaagccttgcctcagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagc 123
OY 241 ggaagcctcagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagc 300
DB 122 ggaagcctcagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagc 63
OY 301 atgacacagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagc 360
DB 62 atgacacagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagcgaagc 4
OY 361 agc 363
DB 3 agc 1

RESULT 4
ABA24907/C
ID ABA24907 standard: DNA: 378 BP.
XX
XX
XX AC ABA24907;
XX
XX DT 23-JAN-2002 (first entry)
XX
XX DE Probe #373 for gene expression analysis in human heart cell sample.
XX
XX DE Human, gene expression; heart; microarray; vascular system; probe;
XX DE cardiovascular disease; hypertension; cardiac arrhythmia;
XX DE congenital heart disease; ss.
XX
XX OS Homo sapiens.
XX
XX PN WO200157274-A2.
XX
XX PD 09-AUG-2001.
XX
XX PF 30-JAN-2001: 2001WO-US00666.
XX
XX PR 04-FEB-2000: 2000US-0180312.
XX PR 26-MAY-2000: 2000US-0207456.
XX PR 30-JUN-2000: 2000US-0608408.
XX PR 03-AUG-2000: 2000US-0632366.
XX PR 21-SEP-2000: 2000US-0234687.
XX PR 27-SEP-2000: 2000US-0236359.
XX PR 04-OCT-2000: 2000GB-0024263.
XX
XX PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX PI Penn SG, Hanzel DK, Chen W, Rank DR:
XX
XX DR WPI: 2001-488899/53.
XX
XX PT Single exon nucleic acid probes for analyzing gene expression in human
XX PT hearts -
XX
XX PS Claim 1: SEQ ID NO 3373: 530pp: English.
XX
XX CC The present invention relates to single exon nucleic acid probes for
XX CC measuring human gene expression in a sample derived from human heart. The
XX CC present sequence is one such probe. The probes may be used for
XX CC predicting, measuring and displaying gene expression in samples derived
XX CC from the human heart via microarrays. By measuring gene expression, the
XX CC probes are useful for predicting, diagnosing, grading, staging,
XX CC monitoring and prognosing diseases of the human heart and vascular system
XX CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and
XX CC congenital heart disease.
XX CC Note: The sequence data for this patent did not form part of the printed
XX CC specification, but was obtained in electronic format directly from WIPO
XX CC at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX SQ Sequence 378 BP; 80 A; 128 C; 97 G; 73 T; 0 other:

Query Match 20.4%; Score 351; DB 22; Length 378;
Best Local Similarity 99.7%; Pred. No. 6.9e-83;
Matches 362; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
DB 1 cctggccctcagatcaagaagcctcttctgtctgtgagcgaaggtgtgagcagc 60
DB 362 cctggccctcagatcaagaagcctcttctgtctgtgagcgaaggtgtgagcagc 303
OY 61 cctctatgagcagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 120
DB 302 cctctatgagcagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 243
OY 121 ggaagatgagcgaagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 180
DB 242 ggaagatgagcgaagcgaagcgaagccttctgtgtgagcgaagcgtgtgagcgaag 183

QY 181 accacaagcttggcttcagagcccaagcccaaggagccagagtgaggaagaatccatcc 240
DB 182 ACCACAAGCTTGGCTTCAGGCCAAGCCCAAGCCAGGGGCGAGGTGAGGAAGTCCATCC 123
QY 241 ggaagctcagatggcagcctgaggaaccctgagggctcaatttcccatctgtgagccgct 300
DB 122 GGAAGTCTGCAATGGCCAGCTGGAGACCTTGGGCTCAATTTCCCATCTGTGAGGCCGCT 63
QY 301 atgaccagctgacaccttccacctccgctactgcatgagccctgtgcatagtgctaaag 360
DB 62 ATGACCAGCTGACACCTTTCACCTCCGCTACTGCTCATGGCCCTGTG-CATAGGTGCTAGGG 4
QY 361 agc 363
DB 3 Agc 1
RESULT 5
AAK03423/C
ID AAK03423 standard; DNA: 378 BP.
XX
AC AAK03423;
DT 05-NOV-2001 (first entry)
DE Human brain expressed single exon probe SEQ ID NO: 3414.
XX
KW Human: brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX
OS Homo sapiens.
XX
PN WO200157275-A2.
XX
PJ 09-AUG-2001.
XX
PJ 30-JAN-2001; 2001WO-US00667.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI: 2001-483446/52.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human
XX brains -
PS Example 4; SEQ ID NO: 3414; 650bp + Sequence Listing: English.
XX
CC The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC brain. They can be used to measure gene expression in brain cell samples,
CC which may enable the diagnosis and improved treatment of nervous system
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC epilepsy and cancers. The present sequence is one of the probes of the
CC invention.
XX
SO Sequence 378 BP; 80 A; 128 C; 97 G; 73 T; 0 other;

Query Match 20.4%; Score 351; DB 22; Length 378;
Best Local Similarity 99.7%; Pred. No. 6,9e-83;
Matches 362; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 ccttgccctcagatcaagaagccttctctctgtgtgcccacaagtgctcgaggcagc 60
DB 362 CTTGGGCCCTCGAGATCAAAAGGCCCTTCTTGTCTGTGGCCAAAGGTGTCCGGCAGC 303
QY 61 cccctcatalggagcagaagaagcagagcttctgtggttgagagagagctgagagtgaa 120
DB 302 CCTCTATGGAGACAGCAAGACAGCTTTGTGGCTGAGAGAGGCTGGCAGGTGAAG 243
QY 121 ggaagatgagagaggtgagagagagatctgtacggttctctgagggctgatalcttgat 180
DB 242 GGAGATGAGAGAGGTAGAGGGGAGATCTGTACGGTTGTTTCTGGGCTCATCTGATAT 183
QY 181 accacaagcttggcttcagagcccaagcccaaggagccagagtgaggaagaatccatcc 240
DB 182 ACCACAAGCTTGGCTTCAGGCCAAGCCCAAGCCAGGGGCGAGGTGAGGAAGTCCATCC 123
QY 241 ggaagctcagatggcagcctgaggaaccctgagggctcaatttcccatctgtgagccgct 300
DB 122 GGAAGTCTGCAATGGCCAGCTGGAGACCTTGGGCTCAATTTCCCATCTGTGAGGCCGCT 63
QY 301 atgaccagctgacaccttccacctccgctactgcatgagccctgtgcatagtgctaaag 360
DB 62 ATGACCAGCTGACACCTTTCACCTCCGCTACTGCTCATGGCCCTGTG-CATAGGTGCTAGGG 4
QY 361 agc 363
DB 3 Agc 1
RESULT 6
AAK28874/C
ID AAK28874 standard; DNA: 378 BP.
XX
AC AAK28874;
DT 06-NOV-2001 (first entry)
DE Human bone marrow expressed single exon probe SEQ ID NO: 3431.
XX
KW Human: bone marrow expressed exon; gene expression analysis; probe;
KW microarray; cancer; leukaemia; lymphoma; myeloma; ss.
XX
OS Homo sapiens.
XX
PN WO200157276-A2.
XX
PJ 09-AUG-2001.
XX
PJ 30-JAN-2001; 2001WO-US00668.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI: 2001-488900/53.
XX
PT Human genome-derived single exon nucleic acid probes useful for
XX analyzing gene expression in human bone marrow -
PS Example 4; SEQ ID NO: 3431; 658bp + Sequence Listing: English.
XX
CC The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC bone marrow. They can be used to measure gene expression in bone marrow
CC samples, which may enable the improved diagnosis and treatment of cancers


```

FT      /note= "Causes R340W"
XX
PN      MO200177305-A2.
PD      18-OCT-2001.
PF      06-APR-2001; 2001WO-SF00765.
XX
PR      07-APR-2000; 2000US-195655P.
XX
PA      (AREX-) AREXIS AB.
PI      Andersson L, Luthman H, Marklund S;
XX
DR      WPI: 2001-657170/75.
P-PSDB: QO847679.
XX
PT      New variants of human AMP-activated protein kinase gamma3 subunit
PT      associated with a metabolic disease e.g. diabetes or obesity and method
PT      for determining a risk estimate of diseases in subject by detecting the
XX      variant -
XX
XX      Disclosure: Fig 5; 25pp: English.
CC
CC      This sequence represents the full length cDNA encoding the human
CC      AMP-activated protein kinase gamma 3 subunit (PRKAG3). Detecting
CC      the presence of the PRKAG3 DNA, or a variant, is useful in determining
CC      a risk estimate of a metabolic disease, such as diabetes or obesity,
CC      in a subject. The variation may occur in exons 3, 4 or 10. In exon
CC      3 variation may be a substitution of a G for a C at nucleotide 320,
CC      resulting in the amino acid substitution P71A; in exon 4 variation may
CC      be a substitution of a T for a C at nucleotide 550; and in exon 10
CC      variation may be a substitution of a T for a C at nucleotide 1037,
CC      resulting in the amino acid substitution R340W. There may also be
CC      nucleotide variation in Intron 6. The numbering of these
CC      variations is based on the full length cDNA as given, rather than on
CC      position 1 of the open reading frame.
XX
SQ      Sequence 1647 BP; 346 A; 502 C; 462 G; 337 T; 0 other;

Query Match          9.8%; Score 168.2; DB 22; Length 1647;
Best Local Similarity 95.6%; Pred No.2.5e-34;
Matches 173; Conservative 0; Mismatches 8; Indels 0; Gaps 0.

QY      1510 ccattccaaacagggtcccctgcgcccccgccctcctcctaaccgaatccaaga 1569
        || | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
DB      1009 cctgcagcatcttcttgctccctgcgcgccgcgccttcctcttaaccgaatccaaga 1068

QY      1570 ttggcgcatcgccacatcccgagacttggtgtgtcgtggagaagaaccatccctgac 1629
        ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
DB      1069 ttggcgcatcgccacatcccgagacttggtgtgtcgtggagaagaaccatccctgac 1128

QY      1630 tgcactgacatctttgtgagccgcggtgtgctgcactgacctgtgtcaagaatgtgg 1689
        ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
DB      1129 tgcactgacatctttgtgagccgcggtgtgctgcactgacctgtgtcaagaatgtgg 1188

QY      1690 t 1690
        |
DB      1189 t 1189

RESULT 12
AAD03296
ID      AAD03296 standard; DNA: 2109 BP.
XX
AC      AAD03296;
XX
DT      13-JUN-2001 (first entry)
XX
DE      Human AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.
XX
XX      Human; gamma subunit; adenosine monophosphate-activated kinase; AMPK;
```

KM	PRKAG3 diabetes: obesity; myopathy; cardiovascular disease; anorectic;
KM	genetic testing; carbohydrate metabolism disorder; skeletal muscle;
KM	cystathione beta synthase; CBS; cardiact; gene therapy; ss.
XX	
OS	Homo sapiens.
FH	
FT	Key Location/Qualifiers
FT	5'UTR 1..471
FT	CDS /tag= a
FT	472..1389
FT	/tag= b
FT	/product= "Human Prkag3 protein"
FT	3'UTR 1390..2109
FT	/tag= c
XX	
XX	WO200120003-A2.
XX	
XX	22-MAR-2001.
XX	
PF	11-SEP-2000; 2000WO-EP09896.
XX	
PR	10-SEP-1999; 99EP-0402236.
PR	18-MAY-2000; 2000EP-0401388.
XX	
PA	(INRG) INRA INST NAT RECH AGRONOMIQUE.
PA	(ANDE/) ANDERSSON L.
PA	(LOOF/) LOOFT C.
PA	(KALM/) KALM E.
XX	
PI	Andersson L, Loof C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
PI	Iannucci N, Gellin J, Le Roy P, Chardon P;
XX	
DR	WPI: 2001-244810/25.
DR	P-PSDB; AAE00221.
XX	
PT	New variants of the gamma subunit of vertebrate adenosine
PT	monophosphate-activated kinase for diagnosis or treatment of disorders
PT	associated with energy metabolism such as diabetes, obesity, and
PT	myopathy -
PS	
PS	Claim 12; Fig 2; 71pp; English.
XX	
CC	The present sequence is a cDNA encoding human adenosine monophosphate
CC	(AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
CC	PRKAG3. Mutation in Prkag3 results in an altered regulation of
CC	carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
CC	useful as therapeutic for treating carbohydrate metabolism disorders such
CC	as diabetes, obesity, and disorders associated with muscle metabolism
CC	activity, and for restoring a normal AMPK function. PRKAG3 sequence
CC	and its functionally altered mutants are useful for the diagnostic
CC	evaluation, genetic testing and prognosis of a metabolic disorder,
CC	preferably a carbohydrate metabolism disorder. Primers that can detect
CC	a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
CC	useful for detecting a dysfunction of carbohydrate metabolism resulting
CC	from the expression of a functionally altered allele of PRKAG3.
CC	Transgenic animal and host cell transformed with PRKAG3 or a
CC	heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC	screening compounds able to modulate AMPK activity. Nucleic acid
CC	encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
CC	in a sequence encoding the first cystathione beta synthase (CBS) domain
CC	of PRKAG3 and is useful in gene therapy.
XX	
XX	
SQ	Sequence 2109 BP; 458 A; 621 C; 560 G; 470 T; 0 other;

Query Match 9.8%; Score 168.2; DB 22; Length 2109;

Best Local Similarity 95.6%; Pred. No. 2.8e-34;

Matches 173; Conservative 0; Mismatches 8; Indels 0; Gaps 0

1510 ccatctcaacagagctccctcgtcgtcccgccctctctctctacgcacatccaaga 1569

|||

909 ccgcgaacattcttgctccctcgtcgtcccgccctctctctctacgcacatccaaga 968

|||||
Db 1035 cgcactggagacatctctcgtggaccgcgctgtgtcctcgcctgcctcgtggtcaccgaaactgg 1094

Search completed: October 3, 2002, 16:30:16
Job time: 14286 sec

GenCore version 4.5
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OM nucleic acid - nucleic search, using sw model

Run on: October 3, 2002, 11:39:20 ; Search time 180.77 Seconds
(without alignments)
2339.884 Million cell updates/sec

Title: US-09-826-581-3

Perfect score: 1722
Sequence: 1 cctggccctcagatcaaga.....gatgagagctcggtcgga 1722

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 38353 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :
1: /cgn2_6/ptodata/2/ina/5A_COMB.seq:*
2: /cgn2_6/ptodata/2/ina/5B_COMB.seq:*
3: /cgn2_6/ptodata/2/ina/5A_COMB.seq:*
4: /cgn2_6/ptodata/2/ina/5B_COMB.seq:*
5: /cgn2_6/ptodata/2/ina/PTODS_COMB.seq:*
6: /cgn2_6/ptodata/2/ina/backfilest1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Length	ID	Description
1	68.8	4.0	US-09-101-146-63	Sequence 63, App1
2	67.6	3.9	US-08-232-463-14	Sequence 14, App1
3	63.8	3.7	US-08-878-989-14	Sequence 14, App1
4	63.8	3.7	US-09-272-796-14	Sequence 14, App1
5	61	3.5	US-08-232-463-14	Sequence 14, App1
6	39.4	2.3	US-09-007-005-17	Sequence 17, App1
7	39.4	2.3	US-09-244-796-17	Sequence 17, App1
8	39	2.3	US-08-658-136-2	Sequence 2, App1
9	39	2.3	US-08-658-136-1	Sequence 1, App1
10	38.6	2.2	US-09-165-264-7	Sequence 7, App1
11	38.6	2.2	US-09-165-264-14	Sequence 14, App1
12	37.4	2.1	US-09-165-264-8	Sequence 8, App1
13	36.2	2.1	US-09-165-264-13	Sequence 13, App1
14	36	2.1	US-09-165-264-11	Sequence 11, App1
15	36	2.1	US-07-959-943-6	Sequence 6, App1
16	36	2.1	US-07-959-943-8	Sequence 8, App1
17	35.8	2.1	US-08-458-568A-11	Sequence 11, App1
18	35.4	2.1	US-09-165-264-12	Sequence 12, App1
19	35.2	2.0	US-09-215-966-21	Sequence 21, App1
20	34.4	2.0	US-08-998-416-876	Sequence 876, App1
21	34.4	2.0	US-09-128-155-16	Sequence 16, App1
22	34.2	2.0	US-08-466-390-3	Sequence 3, App1
23	34.2	2.0	US-08-470-950-3	Sequence 3, App1
24	34.2	2.0	US-08-467-781-3	Sequence 3, App1
25	34.2	2.0	US-08-195-487-3	Sequence 3, App1
26	34.2	2.0	US-08-483-924-3	Sequence 3, App1
27	34.2	2.0	PCT-US93-06160-3	Sequence 3, App1

C 28	34.2	2.0	7705	2	US-08-687-080-115	Sequence 115, App1
C 29	34	2.0	2117	4	US-09-155-768-11	Sequence 11, App1
C 30	34	2.0	6232	4	US-08-456-200B-11	Sequence 11, App1
C 31	34	2.0	7175	1	US-08-455-543A-8	Sequence 8, App1
C 32	34	2.0	7175	2	US-08-193-078B-8	Sequence 8, App1
C 33	34	2.0	7175	2	US-08-223-305C-8	Sequence 8, App1
C 34	34	2.0	7175	2	US-08-149-097D-8	Sequence 8, App1
C 35	34	2.0	7175	3	US-08-949-386-8	Sequence 8, App1
C 36	34	2.0	7175	3	US-08-450-562-8	Sequence 8, App1
C 37	34	2.0	7175	4	US-08-984-709A-8	Sequence 7, App1
C 38	34	2.0	7177	4	US-09-268-163-7	Sequence 1, App1
C 39	34	2.0	7266	3	US-08-713-118-1	Sequence 1, App1
C 40	34	2.0	7266	3	US-09-452-007-1	Sequence 1, App1
C 41	34	2.0	7362	1	US-08-455-543A-7	Sequence 7, App1
C 42	34	2.0	7362	2	US-08-193-078B-7	Sequence 7, App1
C 43	34	2.0	7362	2	US-08-223-305C-7	Sequence 7, App1
C 44	34	2.0	7362	2	US-08-149-097D-7	Sequence 7, App1
C 45	34	2.0	7362	3	US-08-949-386-7	Sequence 7, App1

ALIGNMENTS

RESULT 1
US-09-101-146-63
Sequence 63, Application US/09101146
Patent No. 6124125
GENERAL INFORMATION:
APPLICANT: Dartmouth College, St. Vincent's Institute of
APPLICANT: Medical Research, Kemp et al.
TITLE OF INVENTION: No. 6124125el AMP Activated Protein Kinase
NUMBER OF SEQUENCES: 64
CORRESPONDENCE ADDRESS:
ADDRESSEE: Jane Massey Licata, Esq.
STREET: 66 E. Main Street
CITY: Marlton
STATE: NJ
COUNTRY: USA
ZIP: 08053
COMPUTER READABLE FORM:
MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 MB STORAGE
COMPUTER: IBM PC
OPERATING SYSTEM: WINDOWS 95
SOFTWARE: WORDPERFECT 6.0 FOR WINDOWS
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/101,146
FILING DATE: October 7, 1998
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PN7450
FILING DATE: 8 JAN 1996
ATTORNEY/AGENT INFORMATION:
NAME: Jane Massey Licata
REGISTRATION NUMBER: 32,257
REFERENCE/DOCKET NUMBER: DC-0050
TELECOMMUNICATION INFORMATION:
TELEPHONE: (856) 810-1515
TELEFAX: (856) 810-1454
INFORMATION FOR SEQ. ID NO: 63:
SEQUENCE CHARACTERISTICS:
LENGTH: 1576
TYPE: Nucleic acid
STRANDEDNESS: Single
TOPOLOGY: Linear
ANTI-SENSE: NO
US-09-101-146-63

Query Match 4.0%; Score 68.8; DB 3; Length 1576;
Best Local Similarity 67.4%; Pred. No. 4.2e-09;
Matches 97; Conservative 0; Mismatches 47; Indels 0; Gaps 0;

OY 1240 cagcctgtttgaagctgtctacacccctcagatcaagaacgagatcgcgtctgttct 1299

OY	cctcgagcgcctgacggaaagggaacgtgccctggacatccaaggttgaaggcaagtaccgact	956
Dh	246 ggg	305
OY	957 ccccttgacc	965
Dh	306 agccatgccc	314

RESULT 11
US-09-165-264-14
; Sequence 14, Application US/09165264
; Date of wa 6107510

```

1 APPLICANT: Vinayagamoorthy, Thuralayah
2 TITLE OF INVENTION: Multi-LoCI Genomic Analysis
3 FILE REFERENCE: 44747
4 CURRENT APPLICATION NUMBER: US/09/165,264
5 CURRENT FILING DATE: 1998-10-01
6 NUMBER OF SEQ ID NOS: 14
7 SOFTWARE: PatentIn Ver. 2.1
8 SEQ ID NO: 14
9 LENGTH: 320
10 TYPE: DNA
11 ORGANISM: Artificial Sequence
12 FEATURE:
13 OTHER INFORMATION: Description of Artificial Sequence:Primer sequence
14 US-09-165-264-14

```

Query Match	2.2%;	Score 38.6;	DB 4;	Length 320;
Best Local Similarity	48.8%;	Pred. NO. 0.27;		
Matches 104;	Conservative 0;	Mismatches 109;	Indels 0;	Gaps 0;

[illegible]

```

RESULT 12
US-09-165-264-8
? Sequence 8, Application US/09165264
? Patent No. 6197510
? GENERAL INFORMATION:
? APPLICANT: Vinayagamorthy, Thirulayah
? TITLE OF INVENTION: Multi-Local Genomic Analysis
? FILE REFERENCE: 44/47
? CURRENT APPLICATION NUMBER: US/09/165,264
? CURRENT FILING DATE: 1998-10-01
? NUMBER OF SEQ ID NOS: 14
? SOFTWARE: PatentIn Ver. 2.1
? SEQ ID NO 8
? LENGTH: 319
? TYPE: DNA
? ORGANISM: Artificial Sequence
? FEATURE:
? OTHER INFORMATION: Description of Artificial Sequence:Primer sequence
US-09-165-264-8

```

Query Match	2.28;	Score 37.4;	DB 4;	length 319;
Best Local Similarity	50.38;	Pred. No. 0.56;		
Matches 92;	Conservative 0;	Mismatches 91;	Indels 0;	Gaps 0;

[illegible]

RESULT 13
US-09-165-264-13
; Sequence 13, Application US/09165264
; Patent No. 6107510

```

: APPLICANT: Vinaayagamorthy, Thirulayah
: TITLE OF INVENTION: Multi-Loci Genomic Analysis
: FILE REFERENCE: 44747
: CURRENT APPLICATION NUMBER: US/09/165,264
: CURRENT FILING DATE: 1998-10-01
: NUMBER OF SEQ ID NOS: 14
: SOFTWARE: PatentIn Ver. 2.1
: SEQ ID NO: 13
: LENGTH: 320
: TYPE: DNA
: ORGANISM: Artificial Sequence
: FEATURE:
: OTHER INFORMATION: Description of Artificial Sequence:Primer sequence
US-09-165-264-13

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[illegible]

```

RESULT 14
US-09-165-264-11
Sequence 11, Application US/09165264
Patent No. 6197510
GENERAL INFORMATION:
APPLICANT: Vinayagamoorthy, Thuralayah
TITLE OF INVENTION: Multi-Loc Genomic Analysis
FILE REFERENCE: 44747
CURRENT APPLICATION NUMBER: US/09/165,264
CURRENT FILING DATE: 1998-10-01

```


GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 14:46:08 ; Search time 7316.32 Seconds
(without alignments)
288.886 Million cell updates/sec

Title: US-09-826-581-3_COPY_600_700

Perfect score: 101

Sequence: 1 aggtccccctgtgtgaggaag.....tggtgccctagaagccacag 101

Scoring table:

IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl: *
1: gb_ba: *
2: gb_hlg: *
3: gb_in: *
4: gb_om: *
5: gb_ov: *
6: gb_pal: *
7: gb_ph: *
8: gb_pl: *
9: gb_pr: *
10: gb_ro: *
11: gb_sts: *
12: gb_sy: *
13: gb_un: *
14: gb_vl: *
15: em_ba: *
16: em_fun: *
17: em_hum: *
18: em_in: *
19: em_mu: *
20: em_om: *
21: em_or: *
22: em_ov: *
23: em_pat: *
24: em_ph: *
25: em_pl: *
26: em_ro: *
27: em_sts: *
28: em_un: *
29: em_vl: *
30: em_hlg_hum: *
31: em_hlg_inv: *
32: em_hlg_other: *
33: em_hgo_inv: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	ID	Description
------------	-------	-------	-------	--------	----	-------------

1	101	100.0	1722	6	AX281580	Sequence
2	101	100.0	196554	2	AC073128	Homo sapi
3	101	100.0	206854	9	AC009974	Homo sapi
4	99.4	98.4	152129	2	AC027416	Homo sapi
5	35	34.7	2539	4	U01206	Bos taurus
6	35	34.7	127794	4	AL138688	Human DNA
7	33.2	32.9	692	5	AF41698	Gallus ga
8	33.2	32.9	895	5	S69088	gHox-homoe
9	33.2	32.9	1084	5	CHKPRX1	Gallus gall
10	32.6	32.3	6335	9	D86982	Human mRNA
11	32.6	32.3	54018	2	AC091776	Chlamydom
12	32.6	32.3	65898	2	AC090435	Chlamydom
13	32.6	32.3	161999	2	AL133352	Chlamydom
14	32.6	32.3	172748	2	AC097417	Homo sapi
15	32.6	32.3	203200	9	AC008744	Rattus no
16	32.4	32.1	3761	9	HS8800653	Homo sapi
17	32.4	32.1	4648	9	AF274863	Homo sapi
18	32.4	32.1	5888	9	AF214521	Sus scrof
19	32	31.7	45375	5	AF328738	Agelaius
20	32	31.7	73699	2	AC094322	Rattus no
21	31.8	31.5	104514	2	AC008713	Homo sapi
22	31.8	31.5	135354	2	AC106731	Homo sapi
23	31.8	31.5	136657	2	AL669924	Homo sapi
24	31.8	31.5	221116	2	AC032044	Homo sapi
25	31.6	31.3	171428	9	AC013407	Homo sapi
26	31.6	31.3	175999	2	AC021154	Homo sapi
27	31.6	31.3	183491	9	CNS01D04	Human chr
28	31.4	31.1	63937	2	AC024290	Homo sapi
29	31.4	31.1	67600	2	AC105210	Homo sapi
30	31.4	31.1	88928	2	AC106624	Homo sapi
31	31.4	31.1	157790	2	AL357509	Rattus no
32	31.4	31.1	164959	2	AC087823	Homo sapi
33	31.4	31.1	207636	2	AC019122	Homo sapi
34	31.4	31.1	215692	2	AC090977	Homo sapi
35	31.4	31.1	215727	2	AL358795	Mus muscu
36	31.2	30.9	145556	2	OSJN00092	Homo sapi
37	31.2	30.9	134140	2	AC010758	Oryza sat
38	31.2	30.9	157437	2	AC021669	Homo sapi
39	31.2	30.9	167882	2	AC096852	Sus scrof
40	31.2	30.9	174281	2	AC068338	Homo sapi
41	31.2	30.9	177433	9	AC011747	Homo sapi
42	31.2	30.9	186229	2	AC095022	Sus scrof
43	31.2	30.9	210651	2	AC105137	Homo sapi
44	31	30.7	1105	6	AX090307	Sequence
45	31	30.7	1105	9	AF305687	Homo sapi

ALIGNMENTS

RESULT	1	AX281580	LOCUS	AX281580	DEFINITION	Sequence 3 from Patent WO0177305.	DNA	11linear	PAT	03-NOV-2001
AX281580	AX281580	AX281580	AX281580	AX281580	AX281580	AX281580.1	GI:16608831			

ORGANISM

human.
Homo sapiens
Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi: Mammalia: Eutheria: Primates: Catarrhini: Homnidae: Homo.

REFERENCE

Andersson, L., Luthman, H. and Marklund, S.
Variants of the human amp-activated protein kinase gamma 3 subunit
Patent: WO 0177305-A 3 18-OCT-2001;
Arexis AB (SE)

FEATURES

source

1..1722

/organism="Homo sapiens"

/db_xref="taxon:9606"

BASE COUNT 321 a 504 c 534 g 363 t

ORIGIN

Query Match 100.0%: Score 101: DB 6: Length 1722:
Best Local Similarity 100.0%: Pred. No. 1.3e-16:
Matches 101: Conservative 0: Mismatches 0: Indels 0: Gaps 0:

Oy 1 aggtcccccctgtgaagatgtgagctgggaatctatagcaccagagagggcg99cg 60
|||||
Db 600 AGGTCCTCCCTGTCAGACAGTCGCTGGCAATCTTATGGCACCCAGAGGGCGGGGCGC 659
|||||

Oy 61 gaaggagatcctcctgagagctgtg1gcctagaagcccaag 101
|||||
Db 660 GAGCGAGCTCCTCGAGAGCTGCTGCCCTTAGAGCCACAG 700
|||||

RESULT 2
AC073128/c
LOCUS AC073128 196554 bp DNA linear HTG 21-FEB-2001
DEFINITION Homo sapiens chromosome 2 clone RP11-64705, WORKING DRAFT SEQUENCE,
17 unordered pieces.
AC073128
AC073128.3 GI:13027579
KEYWORDS HTG: HTGS_PHASE1: HTGS_DRAFT: HTGS_FULLTOP.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 196554)
Waterson, R. H.
The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 196554)
Waterson, R. H.
Direct Submission
Submitted (08-JUN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Feb 21, 2001 this sequence version replaced gi:8469048.

COMMENT

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_NH0647005
----- Summary Statistics -----
Sequencing vector: M13: 98%
Sequencing vector: Plasmid: 0%
Chemistry: Dye-terminator Big Dye: 0% of reads
Chemistry: Dye-terminator Big Dye: 0% of reads
Assembly program: Phrap: version 0.990319
Consensus quality: 187795 bases at least Q40
Consensus quality: 190513 bases at least Q30
Insert size: 200000: agarose-fp
Insert size: 194954: sum-of-ctrls
Quality coverage: 5.58 in Q20 bases: sum-of-ctrls
Quality coverage: 5.67 in Q20 bases: sum-of-ctrls

* NOTE: This is a 'working draft' sequence. It currently
* consists of 17 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 1157: contig of 1157 bp in length
* 1158 1257: gap of unknown length
* 1258 3600: contig of 2343 bp in length
* 3601 3700: gap of unknown length
* 3701 5103: contig of 1403 bp in length
* 5104 5203: gap of unknown length
* 5204 8524: contig of 3321 bp in length

* 8525 8624: gap of unknown length
* 8625 11856: contig of 3232 bp in length
* 11857 11956: gap of unknown length
* 11957 15783: contig of 3827 bp in length
* 15784 15883: gap of unknown length
* 15884 21906: contig of 6023 bp in length
* 21907 22005: gap of unknown length
* 22007 28887: contig of 6881 bp in length
* 28888 28987: gap of unknown length
* 28988 35255: contig of 6268 bp in length
* 35256 35355: gap of unknown length
* 35356 44642: contig of 9287 bp in length
* 44643 44742: gap of unknown length
* 44743 58275: contig of 13533 bp in length
* 58276 58375: gap of unknown length
* 58376 73816: contig of 15441 bp in length
* 73817 73916: gap of unknown length
* 73917 92140: contig of 18224 bp in length
* 92141 92240: gap of unknown length
* 92241 113337: contig of 21097 bp in length
* 113338 113437: gap of unknown length
* 113438 130325: contig of 16888 bp in length
* 130326 130425: gap of unknown length
* 130426 149287: contig of 18862 bp in length
* 149288 149388: gap of unknown length
* 149388 196554: contig of 47167 bp in length.

FEATURES
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1. 196554
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="2"
/clone="RP11-64705"
1. 1157
/note="assembly_name:Contig17"
1258. 3600
/note="assembly_name:Contig18"
3701. 5103
/note="assembly_name:Contig19
clone_end:77
vector_side:right"
5204. 8524
/note="assembly_name:Contig20"
8625. 11856
/note="assembly_name:Contig21"
11957. 15783
/note="assembly_name:Contig22"
15884. 21906
/note="assembly_name:Contig23"
22007. 28887
/note="assembly_name:Contig24"
28988. 35255
/note="assembly_name:Contig25"
35356. 44642
/note="assembly_name:Contig26"
44743. 58275
/note="assembly_name:Contig27"
58376. 73816
/note="assembly_name:Contig28"
73917. 92140
/note="assembly_name:Contig29"
92241. 113337
/note="assembly_name:Contig30"
113438. 130325
/note="assembly_name:Contig31"
130426. 149287
/note="assembly_name:Contig32"
149388. 196554
/note="assembly_name:Contig33
clone_end:SP6
vector_side:right"
BASE COUNT 52296 a 46993 c 45889 g 49770 t 1606 others
ORIGIN

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Query Match          100.0%: Score 101: DB 2: Length 196554:
Best Local Similarity 100.0%: Pred. No. 4.4e-17;
Matches 101: Conservative 0: Mismatches 0: Indels 0: Gaps 0:
OY 1 agagccccccgagtgaggaatgaggaatctatgagcaccagaagggcgagggcg 60
    |||
Db 60519 AGGTCCTCTGCTGAGAGAGTGGGCTGGGAATCTTATGGCACCACAGAGGGGGCGG 60460
    |||
OY 61 gaagggaagtcctcctgagcctgagtcgaggaagcccaag 101
    |||
Db 60459 CAGCGAGACTCTCTCTGAGCCTGCTGCTGAGAGAGAGAGAGAGAGAGAGAG 60419
    |||

```

```

RESULT 3          AC009974          206854 bp      DNA      linear      PRI 09-JAN-2002
AC009974/c
LOCUS          AC009974          206854 bp      DNA      linear      PRI 09-JAN-2002
DEFINITION      Homo sapiens BAC clone RP11-459119 from 2, complete sequence.
ACCESSION      AC009974
VERSION        AC009974.9  GI:16799058
KEYWORDS
SOURCE        human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

```

```

REFERENCE
AUTHORS      Sulston,J.E. and Waterston,R.
TITLE        Toward a complete human genome sequence
JOURNAL      Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE      99063792
REFERENCE
AUTHORS      Harris,A. and Cotton,M.
TITLE        The sequence of Homo sapiens BAC clone RP11-459119
JOURNAL      Unpublished (2001)
REFERENCE
AUTHORS      Waterston,R.H.
TITLE        3 (bases 1 to 206854)
JOURNAL      Direct Submission
Submitted (08-SEP-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

```

```

REFERENCE
AUTHORS      4 (bases 1 to 206854)
TITLE        Waterston,R.H.
JOURNAL      Direct Submission
Submitted (08-NOV-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

```

```

REFERENCE
AUTHORS      5 (bases 1 to 206854)
TITLE        Waterston,R.H.
JOURNAL      Direct Submission
Submitted (03-JAN-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

```

```

REFERENCE
AUTHORS      6 (bases 1 to 206854)
TITLE        Waterston,R.
JOURNAL      Direct Submission
Submitted (09-JAN-2002) Department of Genetics, Washington
University 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Nov 8, 2001 this sequence version replaced gi:13431203.

```

```

COMMENT
-----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: saplens@wustl.wustl.edu
-----
Summary Statistics
Center Project name: H_NH0459119
-----

```

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:

all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osogawa,K., Moon,P.Y., Zhao,B., Frengen,E., Tateno,M., Catalanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pletier de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)
VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is RP11-1077K22; the clone sequenced to the right is RP11-64705. Actual start of this clone is at base position 1 of RP11-459119; actual end is at base position 206854 of RP11-459119.

Data from AC079810 and AC073128 was used to finish this clone, AC009974. Polymorphisms have been identified between AC073128 and AC009974. A single plasmid region exists between 38812-38903. An unresolved tandem in the HERV SVA exists between 184390-185163. PCR suggests that approximately 1700 bps are missing.

FEATURES

```

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1. 206854
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/chromosome="2"
/map="2"
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/clone_lib="RPCI-11"
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1..37
/note="match to EST BI059713 (NID:g14467240)"
1..37
/note="match to EST BF183086 (NID:g11061273)"
1..37
/note="match to EST AL567345 (NID:g12920610)"
1..37
/note="match to EST AW880850 (NID:g8042860)"
1..37
/note="match to EST BF304755 (NID:g11251653)"
1..37
/note="similar to Homo sapiens EST B1114348 (NID:g14565249)"
1..37
/note="match to EST BG477625 (NID:g13409904)"
1..37
/note="match to EST BE047599 (NID:g8364652) tz39c01.y1"
3..37
/note="match to EST BE908408 (NID:g10402954)"
4..37
/note="match to EST A1670836 (NID:g4850567) wa04g10.x1"
164..662
/note="match to EST C05773 (NID:g1502549)"
220..221
/note="match to EST BG470047 (NID:g13402322)"
281..344
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281..344
/note="match to EST BE908408 (NID:g10402954)"

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misc_feature	281. .344	/note="match to EST BG477625 (NID:g13409904)"
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misc_feature	281. .344	/note="match to EST BE314060 (NID:g9134719)"
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misc_feature	294. .344	/note="match to EST BG470047 (NID:g13402322)"
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misc_feature	594. .763	/note="match to EST BG470047 (NID:g13402322)"
misc_feature	594. .763	/note="match to EST AM880850 (NID:g8042860)"
misc_feature	594. .731	/note="match to EST BE314060 (NID:g9134719)"
misc_feature	594. .764	/note="match to EST BI059713 (NID:g14467240)"
misc_feature	594. .763	/note="match to EST BG477625 (NID:g13409904)"
misc_feature	594. .742	/note="match to EST BE047599 (NID:g8364652) tz39c01.y1"
misc_feature	594. .763	/note="match to EST BE908408 (NID:g10402954)"
misc_feature	594. .763	/note="match to EST AA043371 (NID:g1521226) zk53e10.r1"
misc_feature	594. .763	/note="match to EST AI670836 (NID:g4850567) wa04g10.x1"
misc_feature	594. .763	/note="match to EST BF183086 (NID:g11061273)"
misc_feature	599. .763	/note="similar to Homo sapiens EST B1114348 (NID:g14565249)"
misc_feature	617. .1084	/note="match to EST AL567345 (NID:g12920610)"
misc_feature	622. .763	/note="match to EST AA481361 (NID:g2210913) zv44e01.r1"
misc_feature	622. .763	/note="match to EST AI860958 (NID:g5514574) w156f05.x1"
misc_feature	622. .763	/note="similar to Mus musculus EST A1196847 (NID:g3749453) u16f01.x1"
misc_feature	684. .763	/note="match to EST BG992568 (NID:g14396638)"
misc_feature	962. .1084	/note="match to EST AI656812 (NID:g4740791) tt54b06.x1"
misc_feature	967. .1084	/note="match to EST BE908408 (NID:g10402954)"
misc_feature	967. .1085	/note="match to EST BF304755 (NID:g11251653)"
misc_feature	967. .1091	/note="match to EST AA043371 (NID:g1521226) zk53e10.r1"
misc_feature	967. .1071	
Query Match	100.0%	Score 101: DB 9: Length 206854;
Best Local Similarity	100.0%	Pred. NO. 4.3e-17;

	Matches	101:	Conservative	0;	Mismatches	0;	Indels	0;	Gaps	0;
Oy	1	aggtcccccctgagtgaaggatggccttgaaattcatgatggaccacaagaaggcgagggcg 60 								
Db	16009	AGGTCCCCCCTGGTAGAGAAGGGCGTGTGGATCTTTATTGGACCCACCAAGAGGCCGC GGCGC 166040 								
Oy	61	gaaggagtcctctctgagacctgacgccctaagaaccaag 101 								
Db	160039	GAGGGAGACTCTCTTGAGAGCTGTGTCCTTGAAAGCCCACC 165999 								
RESULT	4									
LOCUS	AC027416/c									
DEFINITION	Homo sapiens clone RP11-504G11, WORKING DRAFT SEQUENCE, 32 unorderd pieces.									
VERSION	AC027416									
KEYWORDS	AC027416.2 GI:8317289									
SOURCE	HTG; HTGS_PHASEI; HTGS_DRAFT. human.									
ORGANISM	Homo saplens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Euarchonta; Primates; Catarrhini; Homnidae; Homo.									
REFERENCE	Birtten,B., Linton,L., Nusbaum,C. and Landier,E.									
AUTHORS	1 (bases 1 to 152129)									
TITLE	Homo saplens, clone RP11-504G11 unpublished									
JOURNAL REFERENCE	2 (bases 1 to 152129) Birtten,B., Linton,L., Nusbaum,C., Landier,E., Abraham,H., Allen,N., Anderson,S., Baldwin,J., Barra,N., Bastien,V., Beda,F., Boguslavskiy,L., Boukhgalter,B., Brown,A., Burkett,G., Campionavsky,I.L., Bouchkalter,B., Brown,A., Buketki-G., Compianiore,A., Castle,A., Chepel,Y., Colangelo,M., Collins,S., Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,D.S., Dodge,S., Domino,M., Doyle,M., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Glnde,M., Goylette,M., Graham,L., Grand-Pierre,N., Grant,G., Hagos,B., Hearford,A., Horton,L., Howland J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J., LaRoque,K., Lamazes,R., Landers,T., Lehoczyk.J., Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald.P., Marguls.N., McCarthy.H., McEwan,P., McGurk.A., McKernan.K., McPheters.R., Melidrm.T., Menzies,L., Mlhova,T., Miranda,C., Mienga.V., Morrow.J., Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill.D., Oliver,T.M., Olivier,J., Peterson,K., Pierre,N., Pisanic,C., Pollara,V., Raymond.C., Riley.R., Rogov.P., Rothman,D., Roy,A., Santos.R., Schauer,S., Severy,P., Spencer.B., Stange-Thomann,N., Stojanovic.N., Subramanian,A., Talamas,J., Tsfaye-S., Theodore.J., Tirelli,A., Travers,M., Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye.W.J., Young.G., Zainoun,J., Zimmerman,A. and Zody/M.									
COMMENT	All repeats were identified using RepeatMasker: Smit, A.F.A. & Green, P. (1996-1997) http://ftp.genome.washington.edu/RW/RepeatMasker.html									
TITLE	Direct Submission									
JOURNALL	Submitted (30-MAR-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA On Jun 7, 2000 This sequence version replaced gi:7342115.									
COMMENT	Center: Whitehead Institute/ MIT Center for Genome Research Web site: http://www.seq.wimlt.edu Contact: sequence_submissions@genome.wi.mit.edu Project Information Center project name: L7458 Center clone name: 504.G.11 ----- Summary Statistics ----- Sequencing vector: M13; M7815; 100% of reads Chemistry: Dye-terminator Big Dye; 100% of reads Assembly program: Phrap; version 0.960731 Consensus quality: 135376 bases at least Q40 Consensus quality: 143264 bases at least Q30 consensus quality: 146503 bases at least Q20 Insert size: 161000; agarose-fp Insert size: 149029; sum-of-controls									

Quality coverage: 3.1 in Q20 bases; agarose-fp
Quality coverage: 3.3 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 32 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1005: contig of 1005 bp in length
* 1006 1105: gap of 100 bp
* 2402: contig of 1297 bp in length
* 2403 2502: gap of 100 bp
* 3823: contig of 1321 bp in length
* 3824 3923: gap of 100 bp
* 3924 5020: contig of 1097 bp in length
* 5021 5120: gap of 100 bp
* 5121 6161: contig of 1041 bp in length
* 6162 6261: gap of 100 bp
* 6262 7547: contig of 1286 bp in length
* 7548 7647: gap of 100 bp
* 7648 9983: contig of 2336 bp in length
* 9984 10083: gap of 100 bp
* 10084 12556: contig of 2473 bp in length
* 12557 12656: gap of 100 bp
* 12657 15043: contig of 2387 bp in length
* 15044 15143: gap of 100 bp
* 15144 17123: contig of 1980 bp in length
* 17124 17223: gap of 100 bp
* 17224 19466: contig of 2243 bp in length
* 19467 19566: gap of 100 bp
* 19567 21928: contig of 2362 bp in length
* 21929 22028: gap of 100 bp
* 22029 24319: contig of 2291 bp in length
* 24320 24419: gap of 100 bp
* 24420 27059: contig of 2640 bp in length
* 27060 27159: gap of 100 bp
* 27160 30170: contig of 3011 bp in length
* 30171 30270: gap of 100 bp
* 30271 33968: contig of 3698 bp in length
* 33969 34068: gap of 100 bp
* 34069 38179: contig of 4111 bp in length
* 38180 38279: gap of 100 bp
* 38280 42366: contig of 4087 bp in length
* 42367 42466: gap of 100 bp
* 42467 46365: contig of 3899 bp in length
* 46366 46465: gap of 100 bp
* 46466 51285: contig of 4820 bp in length
* 51286 51385: gap of 100 bp
* 51386 55871: contig of 4486 bp in length
* 55872 55971: gap of 100 bp
* 55972 60595: contig of 4624 bp in length
* 60596 60695: gap of 100 bp
* 60696 66595: contig of 5900 bp in length
* 66596 66695: gap of 100 bp
* 66696 73218: contig of 6523 bp in length
* 73219 73318: gap of 100 bp
* 73319 77115: contig of 3797 bp in length
* 77116 77215: gap of 100 bp
* 77216 85022: contig of 7807 bp in length
* 85023 85122: gap of 100 bp
* 85123 93314: contig of 8192 bp in length
* 93315 93414: gap of 100 bp
* 93415 101193: contig of 7779 bp in length
* 101194 101293: gap of 100 bp
* 101294 113090: contig of 11797 bp in length
* 113091 113190: gap of 100 bp
* 113191 123496: contig of 10306 bp in length
* 123497 123596: gap of 100 bp
* 123597 137837: contig of 14241 bp in length
* 137838 137937: gap of 100 bp

FEATURES * 137938 152129: contig of 14192 bp in length.
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1. .152129
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Rp11-504G11"
/clone_1lib="RPC1-11 Human Male BAC"
1. .1005
/note="assembly_fragment"
1106. .2402
/note="assembly_fragment"
2503. .3823
/note="assembly_fragment"
3924. .5020
/note="assembly_fragment"
5121. .6161
/note="assembly_fragment"
6262. .7547
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7648. .9983
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10084. .12556
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12657. .15043
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15144. .17123
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17224. .19466
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vector_slide:left"
19567. .21928
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22029. .24319
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24420. .27059
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27160. .30170
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34069. .38179
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66696. .73218
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73319. .77115
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77216. .85022
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101294. .113090
/note="assembly_fragment"

Query Match 98.4% Score 99.4; DB 2: Length 152129;
Best Local Similarity 99.0%; Pred. No. 1.2e-16;
Matches 100; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 agtcccccctgtgagagtggtcggaatctatgagcaccacgagggcg99g9cg 60

* NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. The true order of the pieces
* is not known and their order in this sequence record is

```

IMPORTANT: this sequence is unfinished and does not necessarily
represent the correct sequence. Work on the sequence is in
progress and the release of this data is based on the understanding
that the sequence may change as work continues. The sequence may
be contaminated with foreign sequence from E.coli, yeast, vector,
phage etc. Order of segments is not known; 800 n's separate
segments. Contig_ID: 00484 Length: 157824bp
Contig_ID: 00808 Length: 15150p
Contig_ID: 01411 Length: 10600p.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 157824: contig of 157824 bp in length
157825 158624: gap of 800 bp

```


Query Match	32.3%	Score 32.6	DB 9	Length 203200
Best Local Similarity	58.9%	Pred. No. 24		
Matches	56	Conservative	0	Mismatches 39; Indels 0; Gaps 0;
QY	7	ccccctgcgagagatgggctcgtgaaactatattatggagaccagagagggcggggcggaaggg	66	
Db	133557	ccccctgcagcgggacatgggctcgtctcttcgaatcggcgctccacagcaggaacttgctc	133498	
QY	67	atctctcctgagaccttgglyccctatagaaccacg	101	
Db	133497	atagcccatgcacgcatgcacgcttgcacgccacg	133463	

Search completed: October 3, 2002, 14:46:57
Job time: 11937 sec

XX		23-JAN-2002	(first entry)	
DT				
XX		Probe #7558 for gene expression analysis in human heart cell sample.		
DE				
XX		Human; gene expression; heart; microarray; vascular system; probe;		
KW		cardiovascular disease; hypertension; cardiac arrhythmia;		
KM		congenital heart disease; ss.		
XX				
OS	Homo sapiens.			
PN	WO200157274-A2.			
PD	09-AUG-2001.			
PF	30-JAN-2001; 2001WO-US00666.			
PR	04-FEB-2000; 2000US-0180312.			
PR	26-MAY-2000; 2000US-0207456.			
PR	30-JUN-2000; 2000US-0608408-			
PR	03-AUG-2000; 2000US-0632366-			
PR	21-SEP-2000; 2000US-0234687.			
PK	27-SEP-2000; 2000US-0236359.			
PA	04-OCT-2000; 2000GB-0024263.			
PI	(MOLE-) MOLECULAR DYNAMICS INC.			
PT	Penn SC, Hanzel DK, Chen W, Rank DR;			
UR	WPI: 2001-488699/53.			
XX				
PT	Single exon nucleic acid probes for analyzing gene expression in human hearts -			
PS	Claim 1: SEQ ID No 7558; 530pp: English.			
CC	The present invention relates to single exon nucleic acid probes for measuring human gene expression in a sample derived from human heart. The present sequence is one such probe. The probes may be used for predicting, measuring and displaying gene expression in samples derived from the human heart via microarrays. By measuring gene expression, the probes are useful for predicting, diagnosing, grading, staging, monitoring and prognosting diseases of the human heart and vascular system e.g. cardiovascular disease, hypertension, cardiac arrhythmias and congenital heart disease.			
CC	Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pcl_sequences.			
SU	Sequence 550 BP; 154 A; 139 C; 127 G; 130 T; 0 other:			
	Query Match	32.3%; Score 32.6; DB 22; Length 550;		
	Best Local Similarity	60.9%; Pred. NO. 4.3;		
	Matches 53; Conservative 0; Mismatches 34; Indels 0; Gaps 0			
OY	10 ctgttgtaggaatctggagccttctaaggacaccagaaggcggcgagagggagt 69 Dd 185 cttactgaagactaaaggccattgttcgccacctacaatagtgycagggctgtggggaag 244			
OY	70 gctccctggagcctgtgcgccttaagaagc 96 Db 245 ccataaatcaaacagctcccctccaac 271			
RESULT	4			
ID	AAK09549			
AC	AAK09549 standard; DNA: 550 BP.			
XX	AAK09549:			
DT	05-NOV-2001 (first entry)			

```

DE Human brain expressed single exon probe SEQ ID NO: 9540.
XX
KW Human; brain expressed exon: gene expression analysis; probe:
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX
OS Homo sapiens.
XX
PN MO200157275-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00667.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI; 2001-483446/52.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human
PT brains -
XX
PS Example 4; SEQ ID NO: 9540; 650pp + Sequence Listing; English.
XX
CC The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC brain. They can be used to measure gene expression in brain cell samples,
CC which may enable the diagnosis and improved treatment of nervous system
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC epilepsy and cancers. The present sequence is one of the probes of the
CC invention.
XX
SO Sequence 550 BP; 154 A; 139 C; 127 G; 130 T; 0 other:

Query Match 32.3%; Score 32.6; DB 22; Length 550;
Best Local Similarity 60.9%; Pred. NO. 4.3;
Matches 53; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

OY 10 ctgtggaagagtggtggtggaatcttatggagaccacagagggcgggcggaaggagat 69
Db 185 ctgatagaagactagcggggggagcaaggtctgtgaccacagatggcgaggggtgctggggaag 244
OY 70 cctcctgagcctgtgtgcctlagaagc 96
Db 245 ccaaaagtaaccagcttcctccaac 271

RESULT 5
AAK35441
ID AAK35441 standard; DNA; 550 BP.
XX
AC AAK35441;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human bone marrow expressed single exon probe SEQ ID NO: 9998.
XX
KW Human; bone marrow expressed exon: gene expression analysis; probe:
KW microarray; cancer; leukaemia; lymphoma; myeloma; ss.
XX
OS Homo sapiens.
XX
PN MO200157276-A2.
XX

```

XX	09-AUG-2001.
XX	
PF	30-JAN-2001; 2001WO-US00668.
XX	
PR	04-FEB-2000; 2000US-0180312.
PR	26-MAY-2000; 2000US-0207456.
PR	30-JUN-2000; 2000US-0608408.
PR	03-AUG-2000; 2000US-0632366.
PR	21-SEP-2000; 2000US-0234687.
PR	27-SEP-2000; 2000US-0236359.
PR	04-OCT-2000; 2000GB-0024263.
XX	
PA	(MOLE-) MOLECULAR DYNAMICS INC.
XX	
PI	Penn SG, Hanzel DK, Chen W, Rank DR;
DR	WPI; 2001-488900/53.
XX	
PT	Human genome-derived single exon nucleic acid probes useful for
XX	analyzing gene expression in human bone marrow -
PS	Example 4; SEQ ID NO: 9998; 658bp + Sequence Listing; English.
CC	The present invention provides a number of single exon nucleic acid
CC	probes which are derived from genomic sequences expressed in the human
CC	bone marrow. They can be used to measure gene expression in bone marrow
CC	samples, which may enable the improved diagnosis and treatment of cancers
CC	such as lymphoma, leukemia and myeloma. The present sequence is one of
XX	the probes of the invention.
SQ	Sequence 550 BP: 154 A; 139 C; 127 G; 130 T; 0 other;
XX	
Query Match	32.3%; Score 32.6; DB 22; Length 550;
Best Local Similarity	60.9%; Pred. No. 4.3;
Matches	53; Conservative 0; Mismatches 34; Indels 0; Gaps 0
OY	10 cctggtcggagcgctggccttggaalctatgaggaccacagaaggcgcgagggagagt 69
Db	185 ctgatgaaagactgcggcgagcaaggctlgtagcacacagatgagcgaggtgctg99gag 244
OY	70 ccttcggagcgctgctgacctagagac 96
Db	245 ccaaaaglaaccagcttccctccaac 271
XX	
RESULT	6
ID	AAL17045
XX	AAL17045 standard; DNA; 550 BP.
AC	AAL17045:
XX	
DT	12-OCT-2001 (first entry)
DE	Probe #6978 for gene expression analysis in human cervical cell sample.
XX	
KM	Probe: human; microarray; gene expression; cervical epithelial cell;
XX	cervical cancer; ss.
OS	Homo sapiens.
XX	
PN	WO200157278-A2.
PD	09-AUG-2001.
XX	
PE	30-JAN-2001; 2001WO-US00670.
XX	
PR	04-FEB-2000; 2000US-0180312.
PR	26-MAY-2000; 2000US-0207456.
PR	30-JUN-2000; 2000US-0608408.
PR	03-AUG-2000; 2000US-0632366.
PR	21-SEP-2000; 2000US-0234687.
PR	27-SEP-2000; 2000US-0236359.
PR	04-OCT-2000; 2000GB-0024263.

PR	27-SEP-2000;	2000US-0236359.
PR	04-OCT-2000;	2000GB-0024263.
XX	(MOLE-)	MOLECULAR DYNAMICS INC.
XX	Penn SG,	Hanzel DK, Chen W, Rank DR;
XX	WPI;	2001-4889901/53.
XX	Human genome-derived single exon nucleic acid probes useful for	
PT	analyzing gene expression in human cervical epithelial cells -	
XX	Claim 25; SEQ ID No 6978;	487bp; English.
XX	The present invention relates to human single exon nucleic acid probes	
CC	(SNP). The present sequence is one such probe. The SNPs are derived	
CC	from human HeLa cells. The SNPs can be used to produce a single exon	
CC	microarray, which can be used for measuring human gene expression in a	
CC	sample derived from human cervical epithelial cells. By measuring gene	
CC	expression, the probes are therefore useful in grading and/or staging	
CC	of diseases of the cervix, notably cervical cancer.	
CC	Note: The sequence data for this patent did not form part of the printed	
CC	specification, but was obtained in electronic format directly from WFO	
CC	at fcp.wipo.int/pub/published_pct_sequences.	
XX	Sequence 550 BP; 154 A; 139 C; 127 G; 130 T; 0 other;	
SQ		
	Query Match	32.3%; Score 32.6; DB 22; Length 550;
	Best Local Similarity	60.9%; Pred. No. 4.3;
	Matches 53; Conservative 0; Mismatches 34; Indels 0; Gaps 0.	
QY	10 ctgttgagagtggtggtggaatctatggtgacccagagggcgagggcgagggagtc 69	
Db	185 ctgttgagagactaggcgggcgaggtgtggtgacccagatggcgaggggtgtctggggagg 244	
QY	70 cctctgtgagcctgtgtgcctctagaagc 96	
Db	245 ccaaaagtaaccagcttcctcccaaac 271	
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ID	AA141154	
XX	AA141154 standard; DNA; 550 BP.	
XX	AA141154;	
DT	17-OCT-2001 (first entry)	
XX	Probe #9840 used to measure gene expression in human placenta sample.	
DE	Probe: microarray; human; placenta; antenatal diagnosis;	
KW	genetic disorder; ss.	
XX	Homo sapiens.	
OS		
XX	MO200157272-A2.	
PN	09-AUG-2001.	
PD		
XX	30-JAN-2001; 2001WO-US00663.	
PF		
XX	04-FEB-2000; 2000US-0180312.	
PR	26-MAY-2000; 2000US-0207456.	
PR	30-JUN-2000; 2000US-0608408.	
PR	03-AUG-2000; 2000US-0632366.	
PR	21-SEP-2000; 2000US-0234687.	
PR	27-SEP-2000; 2000US-0236359.	
PR	04-OCT-2000; 2000GB-0024263.	
XX		
PA	(MOLE-) MOLECULAR DYNAMICS INC.	
XX	Penn SG, Hanzel DK, Chen W, Rank DR;	
XX	WPI;	2001-4889901/53.
XX	Human genome-derived single exon nucleic acid probes useful for	
PT	analyzing gene expression in human cervical epithelial cells -	
XX	Claim 25; SEQ ID No 6978;	487bp; English.
XX	The present invention relates to human single exon nucleic acid probes	
CC	(SNP). The present sequence is one such probe. The SNPs are derived	
CC	from human HeLa cells. The SNPs can be used to produce a single exon	
CC	microarray, which can be used for measuring human gene expression in a	
CC	sample derived from human cervical epithelial cells. By measuring gene	
CC	expression, the probes are therefore useful in grading and/or staging	
CC	of diseases of the cervix, notably cervical cancer.	
CC	Note: The sequence data for this patent did not form part of the printed	
CC	specification, but was obtained in electronic format directly from WFO	
CC	at fcp.wipo.int/pub/published_pct_sequences.	
XX	Sequence 550 BP; 154 A; 139 C; 127 G; 130 T; 0 other;	
SQ		
	Query Match	32.3%; Score 32.6; DB 22; Length 550;
	Best Local Similarity	60.9%; Pred. No. 4.3;
	Matches 53; Conservative 0; Mismatches 34; Indels 0; Gaps 0.	
QY	10 ctgttgagagtggtggtggaatctatggtgacccagagggcgagggcgagggagtc 69	
Db	185 ctgttgagagactaggcgggcgaggtgtggtgacccagatggcgaggggtgtctggggagg 244	
QY	70 cctctgtgagcctgtgtgcctctagaagc 96	
Db	245 ccaaaagtaaccagcttcctcccaaac 271	
RESULT 7		
ID	AA141154	
XX	AA141154 standard; DNA; 550 BP.	
XX	AA141154;	
DT	17-OCT-2001 (first entry)	
XX	Probe #9840 used to measure gene expression in human placenta sample.	
DE	Probe: microarray; human; placenta; antenatal diagnosis;	
KW	genetic disorder; ss.	
XX	Homo sapiens.	
OS		
XX	MO200157272-A2.	
PN	09-AUG-2001.	
PD		
XX	30-JAN-2001; 2001WO-US00663.	
PF		
XX	04-FEB-2000; 2000US-0180312.	
PR	26-MAY-2000; 2000US-0207456.	
PR	30-JUN-2000; 2000US-0608408.	
PR	03-AUG-2000; 2000US-0632366.	
PR	21-SEP-2000; 2000US-0234687.	
PR	27-SEP-2000; 2000US-0236359.	
PR	04-OCT-2000; 2000GB-0024263.	
XX		
PA	(MOLE-) MOLECULAR DYNAMICS INC.	
XX	Penn SG, Hanzel DK, Chen W, Rank DR;	
XX	WPI;	2001-4889901/53.
XX	Human genome-derived single exon nucleic acid probes useful for	
PT	analyzing gene expression in human cervical epithelial cells -	
XX	Claim 25; SEQ ID No 6978;	487bp; English.
XX	The present invention relates to human single exon nucleic acid probes	
CC	(SNP). The present sequence is one such probe. The SNPs are derived	
CC	from human HeLa cells. The SNPs can be used to produce a single exon	
CC	microarray, which can be used for measuring human gene expression in a	
CC	sample derived from human cervical epithelial cells. By measuring gene	
CC		

PR	08-NOV-2000:	2000US-0246525.
PR	08-NOV-2000:	2000US-0246526.
PR	08-NOV-2000:	2000US-0246527.
PR	08-NOV-2000:	2000US-0246528.
PR	08-NOV-2000:	2000US-0246532.
-R	08-NOV-2000:	2000US-0246609.
PR	08-NOV-2000:	2000US-0246610.
PR	08-NOV-2000:	2000US-0246611.
PR	08-NOV-2000:	2000US-0246613.
PR	17-NOV-2000:	2000US-0249207.
PR	17-NOV-2000:	2000US-0249208.
PR	17-NOV-2000:	2000US-0249209.
PR	17-NOV-2000:	2000US-0249210.
PR	17-NOV-2000:	2000US-0249211.
PR	17-NOV-2000:	2000US-0249212.
PR	17-NOV-2000:	2000US-0249213.
PR	17-NOV-2000:	2000US-0249214.
PR	17-NOV-2000:	2000US-0249215.
PR	17-NOV-2000:	2000US-0249216.
PR	17-NOV-2000:	2000US-0249217.
PR	17-NOV-2000:	2000US-0249218.
PR	17-NOV-2000:	2000US-0249244.
PR	17-NOV-2000:	2000US-0249245.
PR	17-NOV-2000:	2000US-0249264.
PR	17-NOV-2000:	2000US-0249265.
PR	17-NOV-2000:	2000US-0249297.
PR	17-NOV-2000:	2000US-0249299.
PR	17-NOV-2000:	2000US-0249300.
PR	01-DEC-2000:	2000US-0250150.
PR	01-DEC-2000:	2000US-0250391.
PR	03-DEC-2000:	2000US-0251030.
PR	05-DEC-2000:	2000US-0251088.
PR	05-DEC-2000:	2000US-0251679.
PR	06-DEC-2000:	2000US-0251479.
PR	08-DEC-2000:	2000US-0251856.
PR	08-DEC-2000:	2000US-0251868.
PR	08-DEC-2000:	2000US-0251869.
PR	08-DEC-2000:	2000US-0251989.
PR	08-DEC-2000:	2000US-0251990.
PR	11-DEC-2000:	2000US-0254097.
PR	05-JAN-2001:	2001US-0259678.
XX	(HUMA-) HUMAN GENOME SCI INC.	
PA	Rosen CA, Barash SC, Ruben SM,	
XI	WPI: 2001-483426/52.	
XX	Nucleic acids encoding human immune/hematopoietic antigen polypeptides,	
PT	useful for preventing, diagnosing and/or treating cancers and	
PT	metastasis -	
XX	Disclosure: SEQ ID NO 31449; 3071pp + Sequence Listing; English.	
PS	AAMK54951 to AAMK64702 encode the human immune/hematopoietic antigen (I)	
CC	amino acid sequences given in AAMK2170 to AAMJ1921. (I) have cytosolic	
CC	activity, and can be used in gene therapy and vaccine production. (I)	
CC	proteins and polynucleotides may be used in the prevention, diagnosis and	
CC	treatment of diseases associated with inappropriate (I) expression. For	
CC	example, they may be used to treat disorders associated with decreased	
CC	expression by rectifying mutations or deletions in a patient's genome	
CC	that affect the activity of (I) by expressing inactive proteins or to	
CC	supplement the patients own production of (I). Additionally, (I)	
CC	polynucleotides may be used to produce the secreted (I), by inserting	
CC	the nucleic acids into a host cell and culturing the cell to express the	
CC	protein. (II) proteins and polynucleotides may be used to prevent,	
CC	diagnose and treat immune/hematopoietic-related diseases, especially	
CC	cancers and cancer metastases of hematopoietic-derived cells. AAMK64703	
CC	to AAMK7694 represent human immune/hematopoietic antigen genomic	
CC	sequences from the present invention. AAK54942 to AAK54950 and AAMK2169	
CC	represent sequences used in the exemplification of the present invention.	
XX	Sequence 680 BP: 116 A; 213 C; 185 G; 166 T; 0 other:	

Query Match	32.3%	Score 32.6	DB 22	Length 680
Best Local Similarity	58.9%	Pred. No. 4.3		
Matches	56	Conservative	0	Mismatches 39; Indels 0; Gaps 0;
Db	629	atagccatgatgcacatgacgcttgcagccacg	663	
Oy	7	ccccctctgtgagagctgtggaatctatgagcaccacagagggcgagggcgagggg	66	
Db	569	ccccctctgtgagagctgtggaatctatgagcaccacagagggcgagggcgagggg	628	
Oy	67	agtcctctgtgagcctgtgtccctagaagccacg	101	
Db	629	atagccatgatgcacatgacgcttgcagccacg	663	
RESULT 9				
AAS81565/C				
ID	AAS81565	standard	CDNA: 3544	BP.
XX	AAS81565;			
XX	AC			
XX	13-FEB-2002	(first entry)		
XX	DNA encoding novel human diagnostic protein #17369.			
XX	Human: chromosome mapping; gene mapping; gene therapy; forensic;			
XX	food supplement; medical imaging; diagnostic; genetic disorder; ss.			
XX	Homo sapiens.			
XX	WO200175067-A2.			
XX	11-OCT-2001.			
XX	30-MAR-2001; 2001WO-US08631.			
XX	31-MAR-2000; 2000US-0540217.			
XX	23-AUG-2000; 2000US-0649167.			
XX	(HYSE-) HYSEQ INC.			
XX	Dmanac RT, Liu C, Tang YT;			
XX	WPI: 2001-639362/73.			
XX	P-PSDB: ABG17378.			
XX	New isolated polynucleotide and encoded polypeptides, useful in			
XX	diagnostics, forensics, gene mapping, identification of mutations			
XX	responsible for genetic disorders or other traits and to assess			
XX	biodiversity			
XX	Claim 1: SEQ ID No 17369; 103pp; English.			
XX	The invention relates to isolated polynucleotide (I) and			
XX	polypeptide (II) sequences. (I) is useful as hybridisation probes,			
XX	polymerase chain reaction (PCR) primers, oligomers, and for chromosome			
XX	and gene mapping, and in recombinant production of (II). The			
XX	polynucleotides are also used in diagnostics as expressed sequence tags			
XX	for identifying expressed genes. (I) is useful in gene therapy techniques			
XX	to restore normal activity of (II) or to treat disease states involving			
XX	(II). (II) is useful for generating antibodies against it, detecting or			
XX	quantitating a polypeptide in tissue, as molecular weight markers and as			
XX	a food supplement. (II) and its binding partners are useful in medical			
XX	imaging of sites expressing (II). (I) and (II) are useful for treating			
XX	disorders involving aberrant protein expression or biological activity.			
XX	The polypeptide and polynucleotide sequences have applications in			
XX	diagnostics, forensics, gene mapping, identification of mutations			
XX	responsible for genetic disorders or other traits to assess biodiversity			
XX	and to produce other types of data and products dependent on DNA and			
XX	amino acid sequences. AAS64197-AAS94564 represent novel human			
XX	diagnostic coding sequences of the invention.			
XX	Note: The sequence data for this patent did not appear in the printed			
XX	note.			

CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 3544 BP; 848 A; 1024 C; 1040 G; 632 T; 0 other;

Query Match 32.3%; Score 32.6; DB 23; Length 3544;
Best Local Similarity 60.9%; Pred. No. 4.7;
Matches 53; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

QY 4 TCCCCCTGTGTAAGAGTGGCTGGAACTLACGGCAACCAGAGGAGGCGGAG 63
DB 2918 TCTGCACTGCGGGCTAGGCGGCTCCAGGCTTGCGCGCCGACGGCTACGCTTGCCCTCAC 2859
QY 64 GAGAGTCTCTCTGAGAGCTGAGCTCCCT 90
DB 2858 GGTCTCTCTCTCTGATCTCTGAACGCT 2832

RESULT 10
AA574057/C
ID AA574057 standard; cDNA: 3750 BP.

XX AA574057;
XX
UT 13-FEB-2002 (first entry)
XX

DE DNA encoding novel human diagnostic protein #9861.

XX Human: chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.

PN WO200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001MO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT.

DR WPI; 2001-639362/73.

DR P-PSDB; ABG09870.

XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity -

PS Claim 1: SEQ ID NO 9861; 103bp; English.

XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. AA564197-AA594364 represent novel human

CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX

SQ Sequence 3750 BP; 841 A; 1043 C; 966 G; 900 T; 0 other;

Query Match 32.1%; Score 32.4; DB 23; Length 3750;
Best Local Similarity 60.0%; Pred. No. 5.4;
Matches 54; Conservative 0; Mismatches 36; Indels 0; Gaps 0;

QY 10 CTGATGAGAGTGGCTGGAACTLACGGCAACCAGAGGAGGCGGAGGAGT 69
DB 475 CTGATGAGAGTGGCTGGAACTLACGGCAACCAGAGGAGGCGGAGGAGT 416
QY 70 CTCCTGAGAGCTGAGCTGAGCTGAGCTGAGCTGAGCTGAGCTGAGCTGAGCT 99
DB 415 CCAAAAGTAACACACTTCTCTCCCAAAAGCA 386

RESULT 11
AAH90107/C
ID AAH90107 standard; cDNA: 3889 BP.

XX AAH90107;
XX

DT 01-OCT-2001 (first entry)
XX

DE Human bone marrow cDNA, SEQ ID NO: 464.

XX Human: bone marrow; anti-inflammatory; cytostatic; neuroprotective;

XX antiviral; antibacterial; antifungal; anti-HIV; haemostatic;

XX immunosuppressive; gene therapy; cytokine cell proliferation;
XX cell differentiation modulator; immune disorder; infection; cancer;

XX human immunodeficiency virus; HIV; autoimmune disorder; haemophilia; ss.

OS Homo sapiens.

PN WO200153453-A2.

PD 26-JUL-2001.

PF 23-DEC-2000; 2000MO-US34960.

PR 21-JAN-2000; 2000US-0488725.

PR 25-APR-2000; 2000US-0552317.

PR 09-JUL-2000; 2000US-0598042.

PR 19-JUL-2000; 2000US-0620312.

PR 03-AUG-2000; 2000US-0653450.

PR 14-SEP-2000; 2000US-0662191.

PR 19-OCT-2000; 2000US-0693036.

PR 30-NOV-2000; 2000US-0250583.

PA (HYSE-) HYSEQ INC.

PI Ford JE, Boyle BJ, Tang YT, Liu C, Asundi V, Chen R, Ma Y;

PI Ren F, Wang J, Werhman T, Xu C, Xue AJ, Yang Y, Zhang J;

PI Zhao QA, Zhou P, Drmanac RT;

DR WPI; 2001-488707/53.
DR P-PSDB; AAM00988.
XX Novel bone-marrow-expressed polynucleotides and polypeptides, useful
XX for treating e.g. cancer and immune deficiency disorders -
XX Claim 1: Page 574-579; 648bp; English.

XX The present sequence is one of 251 novel human polynucleotides
XX expressed in the bone marrow. The polynucleotide and the
XX polypeptide encoded by it are useful in the treatment of various
XX immune deficiencies and disorders. The deficiencies and disorders may
XX be genetic, may be caused by a viral (e.g. HIV), bacterial or fungal

CC infection or may result from an autoimmune disorder, a coagulation
CC disorder (e.g. haemophilia), inhibition of tumour cell proliferation,
CC suppression of an inflammatory response or treatment of a nervous
CC system disorder such as Alzheimer's disease. Detection of the presence
CC or increased expression of the polynucleotide or the protein it
CC encodes is useful for the diagnosis and/or prognosis of one
CC or more types of cancer. The polynucleotide and polypeptide can be
CC used as nutritional sources or supplements and in the screening of
CC chemical compounds as potential drugs.
XX
S0 Sequence 3869 BP; 861 A; 1090 C; 1008 G; 930 T; 0 other;

Query Match	32.1%	Score	32.4	DB	22	Length	3889
Host Local Similarity	60.0%	Pred. No.	5.4				
Matches	54	Conservative	0	Mismatches	36	Indels	0
						Gaps	0

Oy	10	cctggtaggaatcgtgagcaactcttatggacaccagaagaaggcgagagggaggaagt	69
Db	633	CTGTATGAAGACTACGCGGGGGCCAAAGCGTGTGGCACCCAGATGGCAGGGGTGCTGGGAGG	574
Oy	70	cctctcctggagcctggtgcctatagaagccca	99
Db	573	CCAATAATTACCACCTTCCTCCCAAGAACA	544

RESULT 12
AAH90056/C
ID AAH90056 standard; cDNA; 3892 BP.

AC	AAH90056;
XX	
DT	01-OCT-2001 (first entry)
XX	

Human bone marrow cDNA, SEQ ID NO: 300.

KM Human: bone marrow; antineoplastic; cytostatic; neuroprotective;
 KM antiviral; antibacterial; antifungal; anti-HIV; haemostatic;
 KM immunosuppressive; gene therapy; cytokine cell proliferation;
 KM cell differentiation; meningeal disorder; infection; cancer;
 KM human immunodeficiency virus; HIV autoimmune disorder; haemophilia; ss

Homo sapiens.

PN W0200153453-A2.

PD 26-JUL-2001.

PF 23-DEC-2000; 2000WO-US34960.

PR 21-JAN-2000; 2000US-0488725.

PR 09-JUL-2000; 2000US-0598042.

03-AUG-2000; 2000US-0653450.

PR 19-OCT-2000; 2000US-0693036.

NOV 2000 200005-02302635

(MISE) MISCLE INC.

PI Ford JE, Boyle BJ, Tang YT, Liu C, Asundi V, Chen R, Ma Y;
PI Ren F, Wang J, Werhman T, Xu C, Xue AJ, Yang Y, Zhang J;
PI Zhao QA, Zhou B, Drmanac SM.

Zhao QA, Zhou P, Drmanac RT;

DR WPI; 2001-488707/53.

DR P-PSDB; AAM00937.

Novel bone-marrow-expressed polynucleotides and polypeptides, useful for treating e.g. cancer and immune deficiency disorders -

Claim 1; Page 411-412; 648pp; English.

CC The present sequence is one of 251 novel human polynucleotides
CC expressed in the bone marrow. The polynucleotide and the
CC polypeptide encoded by it are useful in the treatment of various
CC immune deficiencies and disorders. The deficiencies and disorders may
CC be genetic, may be caused by a viral (e.g. HIV), bacterial or fungal
CC infection, or may result from an autoimmune disorder, a coagulation
CC disorder (e.g. haemophilia), inhibition of tumour cell proliferation,
CC suppression of an inflammatory response or treatment of a nervous
CC system disorder such as Alzheimer's disease. Detection of the presence
CC or increased expression of the polynucleotide or the protein it
CC encodes is useful for the diagnosis and/or prognosis of one
CC or more types of cancer. The polynucleotide and polypeptide can be
CC used as nutritional sources or supplements and in the screening of
CC chemical compounds as potential drugs.

Query Match	32.1%	Score 32.4	DB 22	Length 3892
Best Local Similarity	60.0%	Pred. No. 5.4		
Matches 54; Conservative	0	Mismatches	36	Indels 0; Gaps 0

OY 10 ctgtgtgaagaaatgttgctctgtagatattcttaggacccacaaagggcgggcgagaaaggaat 63
 Db 633 CTGATGAAGACTAGCCGGGGGCAAGCGCTGTGTGGCCACAGATGGGCGAGGGTGCTGGGAGG 574
 OY 70 cctcctgagacctgtgtccctataagaagccca 99
 Db 573 CCAAAAGTAAACCAAGCTCCCTCCCAAAAACA 544

RESULT	13
AAH89943/c	
ID	AAH89943 standard; cDNA; 4886 BP.

AC AAH89943;

DT 01-OCT-2001 (first entry)

DE Human bone marrow CDNA, SEQ ID NO: 74.

KW Human; bone marrow; antiinflammatory; cytostatic; neuroprotective:

immunosuppressive: gene therapy: cytokine cell proliferation

human immunodeficiency virus: HIV: autoimmune disorder: haemophilia: essential immunodeficiency: immunodeficiency; immune disorder; infection; cancer; cell differentiation

aa
os Homo sapiens.

AA WO200153453-A2
PN

26-JUN-2001

XX
PF 23-DEC-2000. 2000WO-TIS34950XX
PB 31-TAN-3000: 3000115-0488735PR 25-APR-2000; 2000US-0552317.
PB 08-TUL-2000; 2000TC-0508042PR 19-JUL-2000; 2000US-0620312.
03 AUG 2000 05:53:50

PR 14-SEP-2000; 2000US-0662191.

PR 30-NOV-2000; 2000US-0250583.

PA (HYSE-) HYSEQ INC.

PI Ford JE, Boyle BJ, Tang YT, Liu C, Asundi V, Chen R, Ma Y;
PI Ren F, Wang J, Werhman T, Xu C, Xue AJ, Yang Y, Zhang J;
PI Zhao QA, Zhou P, Drmanac RT;
PI

DR WPI; 2001-488707/53.
DR P-PSDB: AAM00824

PT Novel bone-marrow-expressed polynucleotides and polypeptides, useful
PT for treating e.g. cancer and immune deficiency disorders -
XX
XX Claim 1: Page 259-260; 648pp; English.
XX
CC The present sequence is one of 251 novel human polynucleotides
CC expressed in the bone marrow. The polynucleotide and the
CC polypeptide encoded by it are useful in the treatment of various
CC immune deficiencies and disorders. The deficiencies and disorders may
CC be genetic, may be caused by a viral (e.g. HIV), bacterial or fungal
CC infection, or may result from an autoimmune disorder, a coagulation
CC disorder (e.g. haemophilia), inhibition of tumour cell proliferation,
CC suppression of an inflammatory response or treatment of a nervous
CC system disorder such as Alzheimer's disease. Detection of the presence
CC or increased expression of the polynucleotide or the protein it
CC encodes is useful for the diagnosis and/or prognosis of one
CC or more types of cancer. The polynucleotide and polypeptide can be
CC used as nutritional sources or supplements and in the screening of
CC chemical compounds as potential drugs.
XX
XX Sequence 4886 BP; 1117 A; 1370 C; 1266 G; 1133 T; 0 other:
SO

Query Match 32.1%; Score 32.4; DB 22; Length 4886;
Best Local Similarity 60.0%; Pred. No. 5.5;
Matches 54; Conservative 0; Mismatches 36; Indels 0; Gaps 0;

QY 10 ctggtgagagtggtggtggaatcttatggtgcaaccagagcggtggtggtggtggt 69
DB 1616 CTGATGAAACACTAGCGCGGCAAGCCTGTCGACACAGATGCGAGGCTGCGGAGC 1557
QY 70 cctcctgagagcctgtgtgcctagagcca 99
DB 1556 CCAAAAGTAACCACTTCCCTCCAAAGCA 1527
DB

RESULT 14
ABA73753
ID ABA73753 standard; DNA: 231 BP.
XX
XX ABA73753;
XX
XX 01-FEB-2002 (first entry)
XX
XX Human foetal liver single exon nucleic acid probe #22058.
XX
XX Human; foetal liver; gene expression; single exon nucleic acid probe; ss.
XX
XX Homo sapiens.
XX
XX WO200157277-A2.
XX
XX 09-AUG-2001.
XX
XX 30-JAN-2001; 2001WO-US00669.
XX
XX 04-FEB-2000; 2000US-0180312.
XX
XX 26-MAY-2000; 2000US-0207456.
XX
XX 30-JUN-2000; 2000US-0608408.
XX
XX 03-AUG-2000; 2000US-0632366.
XX
XX 21-SEP-2000; 2000US-0234687.
XX
XX 27-SEP-2000; 2000US-0236359.
XX
XX 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX WPI: 2001-483447/52.
XX
XX Human genome-derived single exon nucleic acid probes useful for
XX analyzing gene expression in human foetal liver -
XX

PS Claim 4: SEQ ID NO 22058; 639pp + sequence listing; English.
XX
XX
CC The invention relates to a single exon nucleic acid probe for
CC measuring human gene expression in a sample derived from human foetal
CC liver. The single exon nucleic acid probes may be used for predicting,
CC measuring and displaying gene expression in samples derived from human
CC foetal liver. The present sequence is a single exon nucleic acid
CC probe of the invention.
CC Note: The sequence data for this patent did not form part of the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX Sequence 231 BP; 54 A; 57 C; 63 G; 57 T; 0 other:
SO

Query Match 31.9%; Score 32.2; DB 22; Length 231;
Best Local Similarity 61.2%; Pred. No. 5.3;
Matches 52; Conservative 0; Mismatches 33; Indels 0; Gaps 0;

QY 10 ctggtgagagtggtggtggaatcttatggtgcaaccagagcggtggtggtggtggt 69
DB 146 ctgattgaagactagagcggtggtggtggtggtggtggtggtggtggtggtggt 205
QY 70 cctcctgagagcctgtgtgcctagaa 94
DB 206 ccaaaagtaaccagcttccctccaa 230
DB

RESULT 15
ABA38943
ID ABA38943 standard; DNA: 231 BP.
XX
XX ABA38943;
XX
XX 23-JAN-2002 (first entry)
XX
XX Probe #17409 for gene expression analysis in human heart cell sample.
XX
XX Human; gene expression; heart; microarray; vascular system; probe;
XX cardiovascular disease; hypertension; cardiac arrhythmia;
XX congenital heart disease; ss.
XX
XX Homo sapiens.
XX
XX WO200157274-A2.
XX
XX 09-AUG-2001.
XX
XX 30-JAN-2001; 2001WO-US00666.
XX
XX 04-FEB-2000; 2000US-0180312.
XX
XX 26-MAY-2000; 2000US-0207456.
XX
XX 30-JUN-2000; 2000US-0608408.
XX
XX 03-AUG-2000; 2000US-0632366.
XX
XX 21-SEP-2000; 2000US-0234687.
XX
XX 27-SEP-2000; 2000US-0236359.
XX
XX 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX WPI: 2001-488899/53.
XX
XX Single exon nucleic acid probes for analyzing gene expression in human
XX hearts -
XX
XX Claim 4; SEQ ID NO 17409; 530pp; English.
XX
XX The present invention relates to single exon nucleic acid probes for
XX measuring human gene expression in a sample derived from human heart. The
XX present sequence is one such probe. The probes may be used for
XX predicting, measuring and displaying gene expression in samples derived
XX

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OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:14:46 ; Search time 5701.1 Seconds

(without alignments)
239.110 Million cell updates/sec

Title: us-09-826-581-3_copy_600_700

Sequence: 1 agtccccccgtgtagagag.....tggtagcctagaagccacg 101

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estmu:*
5: em_estcov:*
6: em_estcpl:*
7: em_estro:*
8: em_hic:*
9: qb_est1:*
10: qb_est2:*
11: qb_hic:*
12: qb_gss:*
13: em_gss_hum:*
14: em_gss_inv:*
15: em_gss_pln:*
16: em_gss_vrt:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	36	35.6	682	10	B1960023
2	35.2	34.9	833	10	BE559614
3	35	34.7	377	9	AW15288
4	33	32.7	872	9	AL530561
5	32.8	32.5	656	10	B1960536
6	32.6	32.3	282	9	AA872225
7	32.6	32.3	354	9	AA026388
8	32.6	32.3	539	10	BF808916
9	32.2	31.9	685	9	AL585199
10	32	31.7	735	10	BG309093
11	31.8	31.5	382	10	BE600720
12	31.8	31.5	463	10	BE600139
13	31.8	31.5	474	10	BE598197
14	31.8	31.5	487	10	BE600794
15	31.8	31.5	532	10	BE595429
16	31.6	31.3	695	12	AG083214
17	31.6	31.3	889	10	B1951430

C 18	31.4	31.1	452	9	A1863974
C 19	31.2	30.9	407	9	AU162756
C 20	31.2	30.9	516	9	AU162755
C 21	31.2	30.9	539	9	AL514039
C 22	31.2	30.9	688	9	AL514251
C 23	31.2	30.9	883	12	A2527770
C 24	31	30.7	242	10	BF685935
C 25	31	30.7	469	9	A1087063
C 26	31	30.7	526	10	BE906990
C 27	31	30.7	529	10	BC481234
C 28	31	30.7	531	9	BE019039
C 29	31	30.7	535	10	BE250790
C 30	31	30.7	535	10	BE296883
C 31	31	30.7	537	10	BE294571
C 32	31	30.7	538	10	BE298630
C 33	31	30.7	542	10	BE295054
C 34	31	30.7	553	10	B1160864
C 35	31	30.7	557	10	B1767103
C 36	31	30.7	572	10	BE783728
C 37	31	30.7	581	9	AL538073
C 38	31	30.7	606	9	AW663549
C 39	31	30.7	610	10	BE300397
C 40	31	30.7	611	10	BT254382
C 41	31	30.7	617	10	BE271259
C 42	31	30.7	629	10	BF794131
C 43	31	30.7	640	9	AL531142
C 44	31	30.7	649	9	AL531376
C 45	31	30.7	658	10	BF982575

ALIGNMENTS

RESULT 1
LOCUS B1960023
DEFINITION HVSMEN0022008f Hordeum vulgare rachis EST library HVCDNA0015
(normal) Hordeum vulgare cDNA clone HVSMEN0022008f, mRNA sequence.
ACCESSION B1960023
VERSION B1960023.1 GI:16311278
KEYWORDS EST.
SOURCE
ORGANISM
barley.
Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae ; Triticeae; Hordeum.
1 (bases 1 to 682)
Wing, R., Close, T.J., Kleinof, A., Wise, R., Chin, A., Begum, D., Frisch, D., Atkins, M., Yu, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Gates, R. and Main, D.
Development of a genetically and physically anchored EST resource for barley genomics: Morex rachis cDNA library
Unpublished (2001)
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hg bases = 121
Seq primer: AATTACCTCTACTAAGG
High quality sequence start: 2
High quality sequence stop: 553.
Location/Qualifiers
1..682
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEN0022008f"
/clone_id="Hordeum vulgare rachis EST library HVCDNA0015 (normal)"
/tissue_type="Rachis"

```

/lab_host="TJc121"
/note="Vector: pBluescript SK(-): Site_1: EcoRI; Site_2:
XhoI; Plants were grown at Washington State University,
Pullman, WA in a greenhouse, the rachises were excised and
frozen in liquid nitrogen (Kleinholz lab). In the TJ Close
lab at the University of California, Riverside total RNA
was prepared, poly(A) was purified, one primary
unamplified cDNA library was made, and 1 million pfu were
in vivo excised to give pBluescript SK(-) cDNA phagemids
(Chin). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Rambo, Main). The
sequence has been trimmed to remove vector sequence and
contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders
see Close TJ, Wing R, Kleinholz A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/g99pages/bgn/31/cover.html)"
BASE COUNT      75 a      262 c      174 g      171 t
ORIGIN

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Query Match      35.6% Score 36; DB 10; Length 682;
Best Local Similarity 60.0%; Pred. No. 47;
Matches 60; Conservative 0; Mismatches 40; Indels 0; Gaps 0;

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```

OY 1 aggtcccccgtgtagagtggtggaatcctatgagaccagagggcgagggcg 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 17 ACGCCCCCGCGGTGGAGCGGCGCGCAAGCTGACAGACAGATGCGTGGACG 76
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
OY 61 gaggggagagctcctctgagagctggtgcccctagaagccac 100
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 77 ACGCGTCCCTACTCGCGCGCGCTGCTGCGACCTCTC 116
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

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```

RESULT 2
LOCUS BE559614 833 bp mRNA linear EST 15-AUG-2000
DEFINITION BE559614 601347308F1 NIH_MGC_8 Homo sapiens cDNA clone IMAGE:3688176 5',
mRNA sequence.
ACCESSION BE559614
VERSION BE559614.1 GI:9803323
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

```

```

REFERENCE 1 (bases 1 to 833)
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
TITLES National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabds-femail.nih.gov
Tissue Procurement: Louis M. Staudt, M.D., Ph.D.
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at: image.lnl.gov
Plate: LNCM380 row: P column: 01
High quality sequence stop: 687.
Location/Qualifiers
1. 833

```

```

FEATURES
SOURCE
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3688176"
/clone_lib="NIH_MGC_8"

```

```

/tissue_type="Burkitt lymphoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: Lymph; Vector: pORF7; Site_1: XhoI; Site_2:
EcoRI; cDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCAAGAG(G). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT      215 a      175 c      257 g      186 t
ORIGIN

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```

Query Match      34.9% Score 35.2; DB 10; Length 833;
Best Local Similarity 65.0%; Pred. No. 74;
Matches 52; Conservative 0; Mismatches 28; Indels 0; Gaps 0;

```

```

OY 7 ccctctgtagagagtggtggaatcctatgagaccagagggcgagggcg 66
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 739 CCCCTGCGTTGGATTGGCGGTGCGTCAATGCCGACCTCGCGCGCGCTGGCG 798
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
OY 67 agtctctgtagagctgtg 86
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 799 AAGCGCTGGGCGCGCTGTG 818
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

```

```

RESULT 3
LOCUS AM315288 377 bp mRNA linear EST 25-APR-2001
DEFINITION AM315288 12477 MARC 2BOV Bos taurus cDNA 5', mRNA sequence.
ACCESSION AM315288
VERSION AM315288.1 GI:6744544
KEYWORDS EST.
SOURCE cow.
ORGANISM Bos taurus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovinae; Bos.

```

```

REFERENCE 1 (bases 1 to 377)
AUTHORS Smith,T.P.L., Grosse,W.M., Freking,B.A., Roberts,A.J., Stone,R.T.,
Casas,E., Wray,J.E., White,J., Cho,J., Faehrenkrug,S.C., Bennett
G.L., Heaton,M.P., Laegreid,W.W., Rohrer,G.A., Chitko-McKown,C.G.,
Pietra,G., Holt,I., Karamycheva,S., Liang,F., Quackenbush,J. and
Keeler,J.W.
TITLES Sequence evaluation of four pooled-tissue normalized bovine cDNA
JOURNAL libraries and construction of a gene index for cattle
MEDLINE Genome Res. 11 (4), 626-630 (2001)
COMMENT Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 20
and mismatch 12 options.
PCR PRIMERS
FORWARD: AGGAACACGATGACCAT
BACKWARD: GTTTCACGATCAGCAGC
Plate: 142 row: D column: 8
Seq primer: ATTTACGTCACACTATAG.
Location/Qualifiers
1. 377

```

```

FEATURES
SOURCE
/organism="Bos taurus"
/db_xref="taxon:9913"
/clone_lib="MARC 2BOV"
/tissue_type="pooled"
/lab_host="DH10B"
/note="Vector: PCMV SPORT6; Site_1: XbaI; Site_2: XhoI;
Library made from pooled tissue from testis, thymus,
semitendinosus muscle, longissimus muscle, pancreas,

```


[illegible]

JOURNAL COMMENT	Unpublished (2001) Contact: Frazer Murray Dept. Genomics and Bioinformatics Roslin Institute Roslin, Midlothian, EH25 9PS, UK Tel: +44 (0)131 527 4200 Fax: +44 (0)131 440 0434 Email: frazer.murray@bbsrc.ac.uk Seq primer: T7.
FEATURES	Location/Qualifiers
source	1..685 /organism="Gallus gallus" /db_xref="taxon:9031" /clone="ROS020D01" /clone_lib="BP_Chicken Embryo Library" /tissue_type="Embryo" /dev_stage="5 days old" /lab_host="DH10B" /note="Vector: pBLUESCRIPT SK; Site_1: NotI; site_2: SalI; Cloned unidirectionally. Primer: Oligo dT, 5' adaptor sequence: 5' TCGAACCTCGAG 3'; 3' adaptor sequence: 5' GGGCCCCGCTTTTATTTTTTTTTTTTTTTT 3'."
BASE COUNT	134 a 222 c 224 g 101 t 4 others
ORIGIN	
Query Match	31.9%; Score 32.2; DB 9; Length 685;
Best Local Similarity	65.7%; Pred. No.3.6e+02;
Matches	46: Conservative 0; Mismatches 24; Indels 0; Gaps 0;
Oy	17 ggagtcggacctggaactataggaccacaaggcgcgagggaggtcctctg 76 Db 364 GGAGGAGCGGGCGGCACATGTGTGCCCGCACGAGGAGGCGGNGCTACC CGCGC 423 Oy 77 gagcctgcgtg 86 Db 424 GAGCTTGTTG 433
RESULT 10	
BG309093	735 bp mRNA linear EST 22-OCT-2001
LOCUS	HVSMEC0002B16f Hordeum vulgare seedling shoot EST library
DEFINITION	HVICDNA0003 (/etiolated and unstressed) Hordeum vulgare cDNA clone
ACCESSION	BG309093
VERSION	BG309093.2 GI:16313793
KEYWORDS	EST.
SOURCE	barley.
ORGANISM	Hordeum vulgare Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae ; Triticeae; Hordeum. 1 (bases 1 to 735) Wing,R., Close,T.J., Kleinhofs,A., Wise,R., Begum,D., Frisch,D., Yu .Y., Henry,D., Palmer,M., Rambo,T., Simmons,J., Choi,D.W., Fenton .R.D., Gates,R. and Main,D. Development of a genetically and physically anchored EST resource for barley genomics: Morex unstressed seedling shoot cDNA library Unpublished (2001) On Feb 22, 2001 this sequence version replaced gi:13109940. Contact: Wing RA Clemson University Genomics Institute Clemson University 100 Jordan Hall, Clemson, SC 29634, USA Tel: 864 656 7288 Fax: 864 656 4293 Email: rwing@clemson.edu Total hg bases = 300 Seq primer: AATTACCTCACTAAGCG High quality sequence stop: 718. Location/Qualifiers 1..735
TITLE	
JOURNAL COMMENT	
FEATURES	
source	

/organism="Hordeum vulgare"
 /cultivar="Morex"
 /db_xref="taxon:4513"
 /clone="HVSMEC0002B16f"
 /clone_1lb="Hordeum vulgare seedling shoot EST library
 HVCNDA0003 (Etiolated and unstressed)"
 /tissue_type="Seedling shoot"
 /lab_host="TJC121"

/note="Vector: lambdaZAP. Site 1: EcoRI. Site 2: XhoI;
 Seeds were surface sterilized then germinated under aseptic
 conditions in the dark at room temperature on filter paper
 with water, nystatin and ceftaxime in covered
 crystallization dishes. Five-day old seedling shoots were
 then harvested, total RNA was prepared, poly(A) RNA was
 purified, one primary unamplified cDNA library was made,
 and 1 million pfu were in vivo excised to give phagescript
 SK(-) cDNA phagemids. These steps were performed in the JY
 Close laboratory at the University of California,
 Riverside (Choi, Close, Fenton). Phagemids were plated and
 picked at the Clemson University Genomics Institute (CUGI)
 (Begum, Palmer, Frisch, Atkins and Wang). Plasmid DNA
 preparations, DNA sequencing and sequence analysis were
 performed at CUGI (Wang, Yu, Frisch, Henry, Simmons, Oates
 , Rambo, Main). The sequence has been trimmed to remove
 vector sequence and contains a minimum of 100 bases of
 phred value 20 or above. For more details on library
 preparation and sequence analysis see
 http://www.genome.clemson.edu/projects/barley. To order
 this clone see http://www.genome.clemson.edu/orders Also
 see Close TJ, Wang R, Kiehnhoft A, Wise R (2001)
 Genetically and physically anchored EST resources for
 barley genomics. Barley Genetics Newsletter 31:29-30.
 (http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html)"

BASE COUNT 189 a 156 c 264 g 116 t 10 others
 ORIGIN

Query Match 31.7%: Score 32; DB 10; Length 735;
 Best Local Similarity 62.5%; Pred. No. 4e+02;
 Matches 50; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

OY 10 ctctgagagagctggccttctatgagcaccagagggcgaggcgaggagtc 69
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 559 CTGCTGGCAGCGGCTGGCGGAGGCGGAGGACCGCGCTGAGCGAGGAGT 618
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 OY 70 cctctgagagcctgtgccc 89
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 619 TTGGCTGGCTGTGTGGCCCC 638

RESULT 11 382 bp mRNA linear EST 18-AUG-2000
 BE600720/c
 LOCUS
 DEFINITION P11_91.G10.b1_A002 Pathogen induced 1 (P11) Sorghum bicolor cDNA,
 mRNA sequence.

ACCESSION BE600720
 VERSION BE600720.1 GI:9855905
 KEYWORDS EST.
 SOURCE sorghum.
 ORGANISM Sorghum bicolor

REFERENCE Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
 clade; Panicoideae; Andropogoneae; Sorghum.
 1 (bases 1 to 382)

AUTHORS Cordonnier-Pratt,M.-M., Gingle,A., Dean,R., Sudman,M. and Pratt
 ,L.H.

TITLE An EST database from Sorghum: pathogen-induced plants
 JOURNAL Unpublished (2000)
 COMMENT Contact: Cordonnier-Pratt MM
 Department of Botany
 The University of Georgia
 Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
 Tel: 706 542 1860

Fax: 706 542 1805
 Email: mmp@uga.edu
 Sequences have been trimmed to exclude PolyA, vector and regions
 below phred quality 16. The threshold for highest quality sequence
 is 20.
 Seq primer: JEN REV
 High quality sequence stop: 382
 POLYA-No.

FEATURES
 source location/Qualifiers

1..382
 /organism="Sorghum bicolor"
 /db_xref="taxon:4558"
 /clone_1lb="Pathogen induced 1 (P11)"
 /note="Organ: Anthracnose-infected leaves from
 two-week-old sorghum plants 48 hr after inoculation;
 Vector: phagescript II from lambda Zap II; Site_1: XhoI;
 Site_2: EcoRI. Two-week-old sorghum plants (BRX 623
 cultivar) were infected with pathogen (isolate PRM421 of
 Colletotrichum graminicola, which is a sorghum isolate).
 RNA was prepared from infected leaves harvested from 45
 seedlings 48 hours after inoculation. Note: young
 seedlings (2 weeks old) exhibit juvenile resistant
 reaction, which is an incompatible interaction. As they
 grow older (4 weeks or older), plants resume susceptibility
 to anthracnose disease. The library was made from poly-A
 RNA in the cloning vector lambda ZAP II. Clones to be
 sequenced were prepared by mass excision. WARNING: While
 most or all ESTs are expected to derive from the host
 plant, no effort was made to eliminate ESTs deriving from
 the pathogen."

BASE COUNT 67 a 134 c 90 g 91 t
 ORIGIN

Query Match 31.5%: Score 31.8; DB 10; Length 382;
 Best Local Similarity 59.3%; Pred. No. 4.2e+02;
 Matches 54; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

OY 11 ttgtgagagagtgagccttctatgagcaccagagggcgaggcgaggagtc 70
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 103 TGATGATGTGTGGGCTGTGCTCATGCGGAGCGGAGGAGGAGGAGAG 44
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 OY 71 ctctgagagcctgtgctcctagaagccacg 101
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 43 GGAGAAATGCTGTGTCCGCGCATGCCCATG 13

RESULT 12 463 bp mRNA linear EST 18-AUG-2000
 BE600139/c
 LOCUS
 DEFINITION P11_79.G07.b1_A002 Pathogen induced 1 (P11) Sorghum bicolor cDNA,
 mRNA sequence.

ACCESSION BE600139
 VERSION BE600139.1 GI:9855212
 KEYWORDS EST.
 SOURCE sorghum.
 ORGANISM Sorghum bicolor

REFERENCE Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
 clade; Panicoideae; Andropogoneae; Sorghum.
 1 (bases 1 to 463)

AUTHORS Cordonnier-Pratt,M.-M., Gingle,A., Dean,R., Sudman,M. and Pratt
 ,L.H.

TITLE An EST database from Sorghum: pathogen-induced plants
 JOURNAL Unpublished (2000)
 COMMENT Contact: Cordonnier-Pratt MM
 Department of Botany
 The University of Georgia
 Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
 Tel: 706 542 1860
 Fax: 706 542 1805
 Email: mmp@uga.edu

Sequences have been trimmed to exclude PolyA, vector and regions

below Phred quality 16. The threshold for highest quality sequence is 20.

Seq primer: JEN REV

High quality sequence stop: 462

POLYA=NO.

FEATURES
source Location/Qualifiers

1..463
/organism="Sorghum bicolor"
/db_xref="taxon:4558"
/clone_lib="Pathogen induced 1 (P11)"
/note="Organ: Anthracnose-infected leaves from two-week-old sorghum plants 48 hr after inoculation; Vector: pBluescript II from lambda Zap II; Site 1: XhoI; Site 2: EcoRI; Two-week-old sorghum plants (B7X 623 cultivar) were infected with pathogen (isolate FRM421 of Colletotrichum graminicola, which is a sorghum isolate). RNA was prepared from infected leaves harvested from 45 seedlings 48 hours after inoculation. Note: young seedlings (2 weeks old) exhibit juvenile resistant reaction, which is an incompatible interaction. As they grow older (4 weeks or older), plants resume susceptibility to anthracnose disease. The library was made from poly-A RNA in the cloning vector lambda Zap II. Clones to be sequenced were prepared by mass excision. WARNING: While most or all ESTs are expected to derive from the host plant, no effort was made to eliminate ESTs deriving from the pathogen."

BASE COUNT 79 a 154 c 124 g 106 t
ORIGIN

Query Match 31.5%: Score 31.8: DB 10: Length 463:
Best Local Similarity 59.3%: Pred. No. 4.3e+02:

Matches 54: Conservative 0: Mismatches 37: Indels 0: Gaps 0:

QY 11 tggtagagagtggtggaattctatggtacccagagggcgagggagagtc 70

DB 114 tgcgtatgctgtggcggtgcctcatgcgcagccgacacagcagcagagag 55

QY 71 ctccggagcctgtgtccctagaagccacg 101

DB 54 gcagagatgctgtgtccgccatgccatg 24

RESULT 13 474 bp mRNA linear EST 18-AUG-2000
BE598197/c LOCUS

DEFINITION P11_66_H03.B1_A002 Pathogen induced 1 (P11) Sorghum bicolor cDNA,
mRNA sequence.

ACCESSION BE598197
VERSION BE598197
KEYWORDS GI:9853270
SOURCE EST.

ORGANISM sorghum.

Sorghum bicolor

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC

clade; Panicoidae; Andropogoneae; Sorghum.

REFERENCE 1 (bases 1 to 474)
Cordonier-Pratt,M.-M., Gingle,A., Dean,R., Sudman,M. and Pratt

AUTHORS L.H.

TITLE An EST database from Sorghum: pathogen-induced plants
JOURNAL Unpublished (2000)

COMMENT Contact: Cordonier-Pratt MM
Department of Botany
The University of Georgia
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 542 1805
Email: mmpratt@uga.edu

Sequences have been trimmed to exclude POLYA, vector and regions
below Phred quality 16. The threshold for highest quality sequence
is 20.

Seq primer: JEN REV

High quality sequence stop: 416
POLYA=NO.

FEATURES
source Location/Qualifiers

1..474
/organism="Sorghum bicolor"
/db_xref="taxon:4558"
/clone_lib="Pathogen induced 1 (P11)"
/note="Organ: Anthracnose-infected leaves from two-week-old sorghum plants 48 hr after inoculation; Vector: pBluescript II from lambda Zap II; Site 1: XhoI; Site 2: EcoRI; Two-week-old sorghum plants (B7X 623 cultivar) were infected with pathogen (isolate FRM421 of Colletotrichum graminicola, which is a sorghum isolate). RNA was prepared from infected leaves harvested from 45 seedlings 48 hours after inoculation. Note: young seedlings (2 weeks old) exhibit juvenile resistant reaction, which is an incompatible interaction. As they grow older (4 weeks or older), plants resume susceptibility to anthracnose disease. The library was made from poly-A RNA in the cloning vector lambda Zap II. Clones to be sequenced were prepared by mass excision. WARNING: While most or all ESTs are expected to derive from the host plant, no effort was made to eliminate ESTs deriving from the pathogen."

BASE COUNT 82 a 160 c 126 g 106 t
ORIGIN

Query Match 31.5%: Score 31.8: DB 10: Length 474:
Best Local Similarity 59.3%: Pred. No. 4.3e+02:

Matches 54: Conservative 0: Mismatches 37: Indels 0: Gaps 0:

QY 11 tggtagagagtggtggaattctatggtacccagagggcgagggagagtc 70

DB 105 tgcgtatgctgtggcggtgcctcatgcgcagccgacacagcagcagagag 46

QY 71 ctccggagcctgtgtccctagaagccacg 101

DB 45 gcagagatgctgtgtccgccatgccatg 15

RESULT 14 487 bp mRNA linear EST 18-AUG-2000
BE600794/c LOCUS

DEFINITION P11_90_C08.B1_A002 Pathogen induced 1 (P11) Sorghum bicolor cDNA,
mRNA sequence.

ACCESSION BE600794
VERSION BE600794
KEYWORDS GI:9855769
SOURCE EST.

ORGANISM sorghum.

Sorghum bicolor

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC

clade; Panicoidae; Andropogoneae; Sorghum.

REFERENCE 1 (bases 1 to 487)
Cordonier-Pratt,M.-M., Gingle,A., Dean,R., Sudman,M. and Pratt

AUTHORS L.H.

TITLE An EST database from Sorghum: pathogen-induced plants
JOURNAL Unpublished (2000)

COMMENT Contact: Cordonier-Pratt MM
Department of Botany
The University of Georgia
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 542 1805
Email: mmpratt@uga.edu

Sequences have been trimmed to exclude POLYA, vector and regions
below Phred quality 16. The threshold for highest quality sequence
is 20.

Seq primer: JEN REV

High quality sequence stop: 486

POLYA=NO.

FEATURES Location/Qualifiers

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:18:58 ; Search time 180.77 Seconds
(Without alignments)
137.241 Million cell updates/sec

Title: US-09-826-581-3_COPY_600_700

Perfect score: 101
Sequence: 1 aggtcccccctggtagagag.....tggtagccctagaagccacag 101

Scoring table: IDENTITY_MNC
Gapop 10.0 , Gapext 1.0

Searched: 38353 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 08
Maximum Match 1008
Listing first 45 summaries

Database :
1: /cgn2_6/prodata/2/ina/5A_COMB.seq:*
2: /cgn2_6/prodata/2/ina/5B_COMB.seq:*
3: /cgn2_6/prodata/2/ina/5A_COMB.seq:*
4: /cgn2_6/prodata/2/ina/5B_COMB.seq:*
5: /cgn2_6/prodata/2/ina/PCTUS_COMB.seq:*
6: /cgn2_6/prodata/2/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	27.8	27.5	722	1	US-08-702-344-8
2	27.8	27.5	4411529	4	US-09-103-840A-1
3	27.6	27.3	6822	4	US-09-426-998-3
4	27.6	27.3	7741	4	US-09-426-998-4
5	27.2	26.9	31571	1	US-08-323-443B-1
6	27.2	26.9	53526	3	US-08-658-136-2
7	27.2	26.9	53577	3	US-08-658-136-1
8	26.8	26.5	1190	1	US-08-451-947-9
9	26.8	26.5	1190	2	US-08-424-826A-9
10	26.8	26.5	1190	5	US-08-928-694-9
11	26.8	26.5	1190	5	PCT-US91-06950-9
12	26.8	26.5	3286	4	US-09-211-417-2
13	26.8	26.5	4183	3	US-08-996-083-2
14	26.6	26.3	573	2	US-09-429-516-2
15	26.6	26.3	573	2	US-08-290-665A-140
16	26.6	26.3	573	5	PCT-US95-10398-140
17	26.4	26.1	6763	2	US-08-756-506-23
18	26.4	26.1	10807	1	US-08-206-176-7
19	26.4	25.9	573	2	US-08-756-506-5
20	26.2	25.9	573	5	US-08-290-665A-139
21	26.2	25.9	573	5	PCT-US95-10398-139
22	26.2	25.9	957	4	US-08-836-075A-17
23	26.2	25.9	4403765	4	US-09-103-840A-2
24	26.2	25.7	775	4	US-08-998-416-580
25	25.8	25.5	462	3	US-08-718-388-4
26	25.8	25.5	573	3	US-08-718-388-6
27	25.8	25.5	573	5	US-08-742-923A-1
					Sequence 144, App

C 28	25.8	25.5	1875	4	US-09-422-869-21	Sequence 21, Appl
C 29	25.8	25.5	3247	4	US-08-718-388-4	Sequence 4, Appl
C 30	25.8	25.5	7824	4	US-08-718-388-6	Sequence 6, Appl
C 31	25.8	25.5	16382	4	US-08-718-388-8	Sequence 8, Appl
C 32	25.8	25.5	49136	4	US-09-422-869-1	Sequence 1, Appl
C 33	25.6	25.3	603	4	US-09-134-246-7	Sequence 7, Appl
C 34	25.6	25.3	685	1	US-08-451-947-7	Sequence 7, Appl
C 35	25.6	25.3	685	2	US-08-424-826A-7	Sequence 7, Appl
C 36	25.6	25.3	685	5	US-08-928-694-7	Sequence 7, Appl
C 37	25.6	25.3	685	5	PCT-US91-06950-7	Sequence 7, Appl
C 38	25.6	25.3	11958	4	US-09-134-246-8	Sequence 8, Appl
C 39	25.6	25.3	4403765	4	US-09-103-840A-2	Sequence 2, Appl
C 40	25.6	25.3	4411529	4	US-09-103-840A-1	Sequence 1, Appl
C 41	25.4	25.1	1960	2	US-08-533-306A-1	Sequence 1, Appl
C 42	25.4	25.1	1960	2	US-08-742-923A-1	Sequence 5, Appl
C 43	25.4	25.1	2680	2	US-08-533-306A-5	Sequence 5, Appl
C 44	25.4	25.1	2680	2	US-08-742-923A-5	Sequence 5, Appl
C 45	25.4	25.1	2887	2	US-08-533-306A-3	Sequence 3, Appl

ALIGNMENTS

RESULT 1
US-08-702-344-8
Sequence 8, Application US/08702344
Patent No. 5723315
GENERAL INFORMATION:
APPLICANT: Jacobs, Kenneth
APPLICANT: McCoy, John
APPLICANT: Lavallee, Edward
APPLICANT: Racie, Lisa
APPLICANT: Merberg, David
APPLICANT: Treacy, Maurice
APPLICANT: Spaulding, Vikki
TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
TITLE OF INVENTION: ENCODING THEM
NUMBER OF SEQUENCES: 37
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genetics Institute, Inc.
STREET: 87 Cambridgepark Drive
CITY: Cambridge
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02140
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/702,344
FILING DATE:
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Brown, Scott A.
REGISTRATION NUMBER: 32,724
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8224
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 722 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-08-702-344-8
Query Match 27.5%; Score 27.8; DB 1; Length 722;
Best Local Similarity 62.0%; Pred. No. 10;
Matches 44; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

Query Match	27.38;	Score 27.6;	DB 4;	Length 6822;
Best Local Similarity	56.78;	Pred. NO. 14;		

Sequence 1, Application US/0832443B
Patent No. 5654170

GENERAL INFORMATION:

APPLICANT: KLINGER, KATHERINE W.
APPLICANT: LANDES, GREGORY M.
APPLICANT: BURN, TIMOTHY C.
APPLICANT: CONNORS, TIMOTHY D.
APPLICANT: DACKOWSKI, WILLIAM R.
APPLICANT: GERMINO, GREGORY
APPLICANT: QIAN, PENG

TITLE OF INVENTION: POLYCYSTIC KIDNEY DISEASE GENE

NUMBER OF SEQUENCES: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: Darby & Darby PC
STREET: 805 Third Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10022

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

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COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

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: SOFTWARE: PatentIn Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/323,443B
: FILING DATE: 12-OCT-1994
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: Ludwig, S. Peter
: REGISTRATION NUMBER: 25,351
: REFERENCE/DOCKET NUMBER: 0372/0A462
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (212) 527-7700
: TELEFAX: (212) 753-6237
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 31571 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: HYPOTHEICAL: NO
: ORIGINAL SOURCE:
: ORGANISM: Homo sapiens
: IMMEDIATE SOURCE:
: CLONE: PBD1 GENOMIC
: US-08-323-443B-1

Query Match
Best Local Similarity 58.8%; Score 27.2; DB 1; Length 31571;
Matches 47; Conservative 0; Mismatches 33; Indels 0; Gaps 0;

OY 5 ccccccgtgtagaggtgctgggaatctatgagcaccacagagggcgagggcgaggg 64
Db 16428 CCTGCTGGTCACTCGTGGGAGGCTGACACCTGGGAGAGTAGAGGCCCTGGCAGGAGG 16487
OY 65 ggaatcctcctcgagccttg 84
Db 16488 TGAGGCTCGGGCTCTGGG 16507

RESULT 6
US-08-658-136-2
: Sequence 2, Application US/08658136
: Patent No. 6071717
: GENERAL INFORMATION:
: APPLICANT: KLINGER, KATHERINE W
: APPLICANT: LANDES, GREGORY M
: APPLICANT: BURN, TIMOTHY C
: APPLICANT: CONNORS, TIMOTHY D
: APPLICANT: DACKOWSKI, WILLIAM
: APPLICANT: GERMINO, GREGORY
: APPLICANT: QIAN, FENG
: TITLE OF INVENTION: POLYCYSTIC KIDNEY DISEASE GENE
: NUMBER OF SEQUENCES: 58
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: GENZYME CORPORATION
: STREET: ONE MOUNTAIN ROAD
: CITY: FRAMINGHAM
: STATE: MASSACHUSETTS
: COUNTRY: USA
: ZIP: 01701
: COMPUTER READABLE FORM:
: MEDIUM TYPE: floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/658,136
: FILING DATE:
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: LASSEN, ELIZABETH
: REGISTRATION NUMBER: 31,845

US-08-658-136-1
```

```

: REFERENCE/DOCKET NUMBER: GEN4-17.8
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 508-872-8400
: TELEFAX: 508-872-5415
: INFORMATION FOR SEQ ID NO: 2:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 53526 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: US-08-658-136-2

Query Match
Best Local Similarity 58.8%; Score 27.2; DB 3; Length 53526;
Matches 47; Conservative 0; Mismatches 33; Indels 0; Gaps 0;

OY 5 ccccccgtgtagaggtgctgggaatctatgagcaccacagagggcgagggcgaggg 64
Db 17414 CCTGCTGGTCACTCGTGGGAGGCTGACACCTGGGAGAGTAGAGGCCCTGGCAGGAGG 17473
OY 65 ggaatcctcctcgagccttg 84
Db 17474 TGAGGCTCGGGCTCTGGG 17493

RESULT 7
US-08-658-136-1
: Sequence 1, Application US/08658136
: Patent No. 6071717
: GENERAL INFORMATION:
: APPLICANT: KLINGER, KATHERINE W
: APPLICANT: LANDES, GREGORY M
: APPLICANT: BURN, TIMOTHY C
: APPLICANT: CONNORS, TIMOTHY D
: APPLICANT: DACKOWSKI, WILLIAM
: APPLICANT: GERMINO, GREGORY
: APPLICANT: QIAN, FENG
: TITLE OF INVENTION: POLYCYSTIC KIDNEY DISEASE GENE
: NUMBER OF SEQUENCES: 58
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: GENZYME CORPORATION
: STREET: ONE MOUNTAIN ROAD
: CITY: FRAMINGHAM
: STATE: MASSACHUSETTS
: COUNTRY: USA
: ZIP: 01701
: COMPUTER READABLE FORM:
: MEDIUM TYPE: floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/658,136
: FILING DATE:
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: LASSEN, ELIZABETH
: REGISTRATION NUMBER: 31,845
: REFERENCE/DOCKET NUMBER: GEN4-17.8
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 508-872-8400
: TELEFAX: 508-872-5415
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 53571 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: US-08-658-136-1
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Query Match 26.9%; Score 27.2; DB 3; Length 53577;
Best Local Similarity 58.8%; Pred. No. 23;
Matches 47; Conservative 0; Mismatches 33; Indels 0; Gaps 0;

QY 5 cccccctggtgagagctggcctggaattctatggcaccacagagggcgaggcgagg 64
DB 17413 CCTCTCGGCTACTGCTGGGAGGCTGACACTGGGAGAGTACAGGCCCGTGGCAGGAG 17472

OY 65 ggaagtcctctggagcctg 84
DB 17473 TGAGCCCTCGGCTCGCTCGG 17492

RESULT 8
US-08-451-947-9/c
Sequence 9, Application US/08451947
Patent No. 5702906
GENERAL INFORMATION:
APPLICANT: GENE TECH, INC.
APPLICANT: ROSENTHAL, ARNON
TITLE OF INVENTION: NOVEL NEUTROTROPIC FACTOR
NUMBER OF SEQUENCES: 100
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genentech, Inc.
STREET: 460 Point San Bruno Blvd
CITY: South San Francisco
STATE: California
COUNTRY: USA
ZIP: 94080
COMPUTER READABLE FORM:
MEDIUM TYPE: 5.25 inch, 360 Kb floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: patin (Genentech)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/451.947
FILING DATE:
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/426419
FILING DATE: 19-APR-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/030013
FILING DATE: 22-MAR-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/648482
FILING DATE: 31-JAN
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/587707
FILING DATE: 1991
ATTORNEY/AGENT INFORMATION:
NAME: Torchia, Timothy E.
REGISTRATION NUMBER: 36,700
REFERENCE/DOCKET NUMBER: 666P2C102
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415/225-8674
TELEFAX: 415/952-9881
TELEX: 910/371-7168
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1190 bases
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: Linear
US-08-451-947-9

Query Match 26.5%; Score 26.8; DB 1; Length 1190;
Best Local Similarity 57.0%; Pred. No. 20;
Matches 49; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

OY 11 tggtagagagtggtggaattctatggcaccacagagggcgaggcgaggc 70
DB 469 TGTCGAGATGGGCTTAGACATCGACACTGGGAGAGGAAAAATGAGGGGATG 410

DB 469 TGTCGAGATGGGCTTAGACATCGACACTGGGAGAGGAAAAATGAGGGGATG 410

QY 71 ctctggagcctgggacctagaagc 96
DB 409 CGAGGAGCCTGGGGAGCAGGAGC 384

RESULT 9
US-08-424-826A-9/c
Sequence 9, Application US/08424826A
Patent No. 5830858
GENERAL INFORMATION:
APPLICANT: Rosenthal, Arnon
TITLE OF INVENTION: NOVEL NEUTROTROPIC FACTOR
NUMBER OF SEQUENCES: 98
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genentech, Inc.
STREET: 460 Point San Bruno Blvd
CITY: South San Francisco
STATE: California
COUNTRY: USA
ZIP: 94080
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Winpatin (Genentech)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/424.826A
FILING DATE: 19-APR-1995
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/240387
FILING DATE: 10-MAY-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/648482
FILING DATE: 31-JAN-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/587707
FILING DATE: 25-SEP-1990
ATTORNEY/AGENT INFORMATION:
NAME: Torchia, Ph.D., Timothy E.
REGISTRATION NUMBER: 36,700
REFERENCE/DOCKET NUMBER: P0666PIC2
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415/225-8674
TELEFAX: 415/952-9881
TELEX: 910/371-7168
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1190 base pairs
TYPE: Nucleic Acid
STRANDEDNESS: Single
TOPOLOGY: Linear
US-08-424-826A-9

Query Match 26.5%; Score 26.8; DB 2; Length 1190;
Best Local Similarity 57.0%; Pred. No. 20;
Matches 49; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

QY 11 tggtagagagtggtggaattctatggcaccacagagggcgaggcgaggc 70
DB 469 TGTCGAGATGGGCTTAGACATCGACACTGGGAGAGGAAAAATGAGGGGATG 410

OY 71 ctctggagcctgggacctagaagc 96
DB 409 CGAGGAGCCTGGGGAGCAGGAGC 384

RESULT 10
US-08-928-694-9/c

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Sequence 9, Application US/08928694
Patent No. 6037320
GENERAL INFORMATION:
APPLICANT: ROSENTHAL, ARNON
TITLE OF INVENTION: NOVEL NEUROTROPHIC FACTOR
NUMBER OF SEQUENCES: 100
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genentech, Inc.
STREET: 1 DNA Way
CITY: South San Francisco
STATE: California
COUNTRY: USA
ZIP: 94080
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Minipain (Genentech)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/928,694
FILING DATE: 12-Sep-1997
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/451947
FILING DATE: 26-MAY-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/426419
FILING DATE: 19-APR-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/030013
FILING DATE: 22-MAR-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/648482
FILING DATE: 31-JAN
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/587707
FILING DATE: 1991
ATTORNEY/AGENT INFORMATION:
NAME: Torchia, Phd., Timothy E.
REGISTRATION NUMBER: 36,700
REFERENCE/DOCKET NUMBER: P0666P2C1D2C1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650/225-8674
TELEFAX: 650/952-9881
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1190 base pairs
TYPE: Nucleic Acid
STRANDEDNESS: Single
TOPOLOGY: Linear
US-08-928-694-9

Query Match 26.5%; Score 26.8; DB 3; Length 1190;
Best Local Similarity 57.0%; Pred. No. 20;
Matches 49; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

OY 11 tggtagagagctgggagcttgaagcaccagagggcggggagggagagtc 70
    |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
DB 469 TGTCGAGATGGGTTAGGACTCCAAATGACACTGGGAGAGGAAATGAGGGGATG 410
OY 71 ctctggagcctgtgcccctagaagc 96
    |||||||  |||||||  |||||||
DB 409 CGAGGAGGCGCTGGGAGGAGCGAGC 384

RESULT 11
PCT-US91-06950-9/c
Sequence 9, Application PC/TUS9106950
GENERAL INFORMATION:
APPLICANT: GENENTECH, INC.
APPLICANT: ROSENTHAL, ARNON
TITLE OF INVENTION: NOVEL NEUROTROPHIC FACTOR
```

```
NUMBER OF SEQUENCES: 100
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genentech, Inc.
STREET: 460 Point San Bruno Blvd
CITY: South San Francisco
STATE: California
COUNTRY: USA
ZIP: 94080
COMPUTER READABLE FORM:
MEDIUM TYPE: 5.25 inch, 360 Kb floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: pain (Genentech)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US91/06950
FILING DATE: 19910924
CLASSIFICATION: 436
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/648482
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/587707
ATTORNEY/AGENT INFORMATION:
NAME: Hensley, Max D.
REGISTRATION NUMBER: 27,043
REFERENCE/DOCKET NUMBER: 666P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415/266-1994
TELEFAX: 415/952-9881
TELEX: 910/371-7168
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1190 bases
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
PCT-US91-06950-9

Query Match 26.5%; Score 26.8; DB 5; Length 1190;
Best Local Similarity 57.0%; Pred. No. 20;
Matches 49; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

OY 11 tggtagagagctgggagcttgaagcaccagagggcggggagggagagtc 70
    |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
DB 469 TGTCGAGATGGGTTAGGACTCCAAATGACACTGGGAGAGGAAATGAGGGGATG 410
OY 71 ctctggagcctgtgcccctagaagc 96
    |||||||  |||||||  |||||||
DB 409 CGAGGAGGCGCTGGGAGGAGCGAGC 384

RESULT 12
US-09-211-417-2/c
Sequence 2, Application US/09211417A
Patent No. 6177254
GENERAL INFORMATION:
APPLICANT: Rattner, Jerome B
APPLICANT: Whitehead, Clark M
TITLE OF INVENTION: NUCLEOLUS AUTOANTIGENIC MARKER FOR SYSTEMIC LUPUS
TITLE OF INVENTION: ERHHEMATOSUS
TITLE OF INVENTION: GenBank
FILE REFERENCE: UCC1
CURRENT APPLICATION NUMBER: US/09/211,417A
CURRENT FILING DATE: 1998-12-15
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 2
LENGTH: 3286
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: Nucleic Acid Sequence of ASE-1
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```

: SOFTWARE: WORDPERFECT 5.1
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/290,665A
: FILING DATE: 15-AUG-1994
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: RICHARD W. BORK
: REGISTRATION NUMBER: 36,459
: REFERENCE/DOCKET NUMBER: 2026-4116
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (212) 751-4800
: TELEFAX: (212) 751-6849
: TELEX: 421792
: INFORMATION FOR SEQ ID NO: 140:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 573 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: ORGANISM: homosapiens
: INDIVIDUAL ISOLATE: 28
: US-08-290-665A-140

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Query Match      26.3%; Score 26.6; DB 2; Length 573;
Best Local Similarity 56.2%; Pred. No. 21;
Matches 50; Conservative 0; Mismatches 39; Indels 0; Gaps 0;

QY 5 cccccctgtaggagctggcctggaatctatgggcacccagaggcgcgagcgaag 64
   ||| | | | | | | | | | | | | | | | | | | | | | | | | |
Db 359 CCCAATTGCGGACCTCCGCGGGATCATTTGGGCCCCAGACGTCGAGAGCGCGCG 300
   ||| | | | | | | | | | | | | | | | | | | | | | | | | |
QY 65 ggaatcctcctgagagcctgtagcctaga 93
   || | | | | | | | | | | | | | | | | | | | | | | | | |
Db 299 GCGGACAGAGAGCCCAACTGCCCCACCACA 271
   || | | | | | | | | | | | | | | | | | | | | | | | | |

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Search completed: October 3, 2002, 16:21:45
Job time: 16945 sec


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PN      WO200177305-A2.
XX      18-OCT-2001.
PD      06-APR-2001: 2001WO-SE00765.
XX      07-APR-2000: 2000US-19565P.
XX      (AREX-) AREXIS AB.
XX      Andersson L, Luthman H, Marklund S;
XX      WPI: 2001-657170/75.
DR      P-PSDB: Q0847679.
XX      New variants of human AMP-activated protein kinase gamma3 subunit
PT      associated with a metabolic disease e.g. diabetes or obesity and method
PT      for determining a risk estimate of diseases in subject by detecting the
PT      variant.
XX      Disclosure: Fig 5; 25pp: English.
XX      This sequence represents the full length cDNA encoding the human
CC      AMP-activated protein kinase gamma 3 subunit (PRKAG3). Detecting
CC      the presence of the PRKAG3 DNA, or a variant, is useful in determining
CC      a risk estimate of a metabolic disease, such as diabetes or obesity,
CC      in a subject. The variation may occur in exons 3, 4 or 10. In exon
CC      3 variation may be a substitution of a G for a C at nucleotide 320,
CC      resulting in the amino acid substitution P71A; in exon 4 variation may
CC      be a substitution of a T for a C at nucleotide 550; and in exon 10
CC      variation may be a substitution of a T for a C at nucleotide 1037,
CC      resulting in the amino acid substitution R340W. There may also be
CC      nucleotide variation in intron 6. The numbering of these
CC      variations is based on the full length cDNA as given, rather than on
CC      position 1 of the open reading frame.
XX      Sequence 1647 BP: 346 A: 502 C: 462 G: 337 T: 0 other:
SO

Query Match      100.0%; Score 1647; DB 22; Length 1647;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 1647; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      1 ttggtctggggtgtggtccacatgagacggcgtgagacagcactgagcagagcccttc 60
DB      1 ttggtctggggtgtggtccacatgagacggcgtgagacagcactgagcagagcccttc 60
OY      61 ctggagcaagccttgggtctgagcatcaagaagtgtggtcttccttagagcaagaacag 120
DB      61 ctggagcaagccttgggtctgagcatcaagaagtgtggtcttccttagagcaagaacag 120
OY      121 cagctcatggtcattacacagcgtgtgacagcagctcagaaagaatccgttgggaaacgag 180
DB      121 cagctcatggtcattacacagcgtgtgacagcagctcagaaagaatccgttgggaaacgag 180
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DB      181 ggcctaaacgttgaatgagacaaagcagaatcgggtgaggaaggagagccacagttca 240
OY      241 gggggaaggtcccggtccagagcagctgagtcacacggcgtggagggccacattccc 300
DB      241 gggggaaggtcccggtccagagcagctgagtcacacggcgtggagggccacattccc 300
OY      301 caagacacaccccttggctcaagctgatactgcgagggtgggacactccaacaacaggtg 360
DB      301 caagacacaccccttggctcaagctgatactgcgagggtgggacactccaacaacaggtg 360
OY      361 ggaactgctccctctgagcttgaagcctcagcagcgtccacagcaagaatgattgga 420
DB      361 ggaactgctccctctgagcttgaagcctcagcagcgtccacagcaagaatgattgga 420
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DB      421 gctggcacaaggagttcccaagcacaagagcctggaggtgagcttaagaagcctgcgga 480

DB      421 gctggcacaaggagttcccaagcacaagagcctggaggtgagcttaagaagcctgcgga 480
OY      481 agagaagcctgctgtgctgtcccgcaaggccattcccaactggtcgtgagatga 540
DB      481 agagaagcctgctgtgctgtcccgcaaggccattcccaactggtcgtgagatga 540
OY      541 cgaactgagaaacccggcgccagatctacatgctgtcattatgagagacacgtgta 600
DB      541 cgaactgagaaacccggcgccagatctacatgctgtcattatgagagacacgtgta 600
OY      601 cgaatgcattgagcaactagctccaagctagatcattctgcacacatgctgagatcaaga 660
DB      601 cgaatgcattgagcaactagctccaagctagatcattctgcacacatgctgagatcaaga 660
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DB      661 ggcctctctgtctgtgagcacaagctgtgaggaagccctctatgagacagaaga 720
OY      721 gcaagagcttggggagatgctgacatcaactgaacttacctggtgctgcatcgtacta 780
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DB      961 gtcaagcaacgtactccaatccatccacaacaagaagcctgtcctgacatctt 1020
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DB      1021 tggctccctgtgctcccgccctctctccctacacatccaaagtctgggacatctt 1080
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DB      1081 cacattccgagaactgtgtgtgtgtgtgagacagaccacttcctgactgacaggaat 1140
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DB      1141 ctgttgagaccggcgtgtgtctgacactgacctgtgtgcaagaatgtgtgacgtgcgtgg 1200
OY      1201 cctctatcccgctttgattgattcaactgtgctgcccagaacacttaacaacacttga 1260
DB      1201 cctctatcccgctttgattgattcaactgtgctgcccagaacacttaacaacacttga 1260
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DB      1321 ccaagcccacagaagccttgggggaagtgtatcgacagaagattgtctcgagggtlaacag 1380
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DB      1441 tcaaggcactgtgtctcagccctgtgcatcagatgacctgtggggcctggaagaatctgag 1500
OY      1501 tctctcaatcccaagccaactgtgacaaccttggaagccaatggaagggaactcagc 1560
DB      1501 tctctcaatcccaagccaactgtgacaaccttggaagccaatggaagggaactcagc 1560
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OY 1621 ggcacatgacacagcctcttaagcttc 1647
DB 1621 ggcacatgacacacagcctcttaagcttc 1647

RESULT 2
AAD03320
ID AAD03320 standard; cDNA: 2115 BP.
XX
AC AAD03320;
XX
DT 13-JUN-2001 (first entry)
XX
DE Human AMPK gamma subunit muscle-specific isoform, complete PRKAG3 cDNA.
XX
KW Human; gamma subunit; adenosine monophosphate-activated kinase; AMPK;
KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
KW cystathione beta synthase; CBS; cardiact; gene therapy; ss.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT CDS 1..1395
FT FT /*tag= a
FT FT /product= "Human complete Prkag3 protein"
XX
PN WO200120003-A2.
XX
PD 22-MAR-2001.
XX
PF 11-SEP-2000: 2000WO-EP09896.
XX
PR 10-SEP-1999: 99EP-0402336.
PR 18-MAY-2000: 2000EP-0401388.
XX
PA (INRG ) INRA INST NAT RECH AGRONOMIQUE.
PA (ANDE/) ANDERSSON L.
PA (LOOF/) LOOFT C.
PA (KALM/) KALM E.
XX
PI Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
PI Tamnucellil N, Gellin J, Le Roy P, Chardon P;
DR MPI: 2001-244810/25.
DR P-PSDB: AME00223.
XX
XX New variants of the gamma subunit of vertebrate adenosine
PT monophosphate-activated kinase for diagnosis or treatment of disorders
PT associated with energy metabolism such as diabetes, obesity, and
PT myopathy -
XX
PS Claim 12: Page 65-68: 71pp: English.
XX
XX The present sequence is a cDNA encoding human adenosine monophosphate
CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
CC complete PRKAG3. Mutation in Prkag3 results in an altered regulation of
CC carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
CC useful as therapeutic for treating carbohydrate metabolism disorders such
CC as diabetes, obesity, and disorders associated with muscle metabolism
CC such as myopathy and cardiovascular diseases, to modulate AMPK
CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
CC and its functionally altered mutants are useful for the diagnostic
CC evaluation, genetic testing and prognosis of a metabolic disorder,
CC preferably a carbohydrate metabolism disorder. Primers that can detect
CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
CC useful for detecting a dysfunction of carbohydrate metabolism resulting
CC from the expression of a functionally altered allele of PRKAG3.
```

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CC Transgenic animal and host cell transformed with PRKAG3 or a
CC heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC screening compounds able to modulate AMPK activity. Nucleic acid
CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
CC in a sequence encoding the first cystathione beta synthase (CBS) domain
CC of PRKAG3 and is useful in gene therapy.
XX
SQ Sequence 2115 BP; 460 A; 622 C; 562 G; 471 T; 0 other;

Query Match 88.2%; Score 1453; DB 22; Length 2115;
Best Local Similarity 97.3%; Pred. No. 0;
Matches 1501; Conservative 0; Mismatches 35; Indels 7; Gaps 2;

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DB 1 atgagcttccttagagcaagaacacacagcattatggtccatccacgctgtgacagcagc 60
OY 155 tcagaaagaatccgttgggaacgagggccaaagccttgaatgagacagagcagaagctg 214
DB 61 tcagaaagaatccgttgggaacgagggccaaagccttgaatgagacagagcagaagctg 120
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DB 181 tcaccgggcttgaggccacattccccaagacacacaccttggctcctaaagtatctgccc 240
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DB 241 ggggtgggcactccaccaaagggtgggactgctccctctgactgtacagctcagct 300
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DB 301 gcaagctccagacagatgatgtgagctggtgccaaggaggttccccaagacagagcttgg 360
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DB 361 gagtgtgagtagaaggcctgtctggaagaagagccttgccctgtgcttccccaaggagcc 420
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DB 421 ccatttcccaagctggcgtgggatacgaactggtggaaacccggcgccagatctcaatg 480
OY 575 cgcttcattcagagacacacccgtcactgactgacatggaactagctccaagctatgctc 634
DB 481 cgcttcattcagagacacacccgtcactgactgacatggaactagctccaagctatgctc 540
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DB 841 catgcgcctgtctctgtgacccggtgtcaggaacgtaactccacatcctacacacaaa 900
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Db 1021 gcaccacatcctgaactgcacatctcttgaggacgcgcgtgtgtctgcactgccttg 1080
Oy 1175 gtcaacgaatgtgtcagtgctgtggcctctatccgcctcttgatgtatccactgtgc 1234
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RESULT 3
AAD03296
ID AAD03296 standard; DNA: 2109 BP.
XX
AC AAD03296;
XX
DT 13-JUN-2001 (first entry)
XX
DE Human AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.
XX
KW Human: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
KW cystathione beta synthase; CBS; cardiant; gene therapy; ss.
XX
OS Homo sapiens.
XX
XX Key Location/Qualifiers
XX FH 1..471
XX FT 5'UTR
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XX FT 3'UTR 1390..2109
XX FT /*tag= c
XX PN W0200120003-A2.

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XX
PD 22-MAR-2001.
XX
PF 11-SEP-2000; 2000WO-EP09896.
XX
PR 10-SEP-1999; 99EP-0402236.
PR 18-MAY-2000; 2000EP-0401388.
XX
PA (INRG ) INRA INST NAT RECH AGRONOMIQUE.
PA (ANDE/) ANDERSSON L.
PA (LOOF/) LOOFT C.
PA (KALM/) KALM E.
XX
PI Andersson L, Loof C, Kalm E, Milan D, Robic A, Rogel-Gallard C;
PI Iannuccelli N, Gellin J, Le Roy P, Chardon P.
XX
DR MPI: 2001-244810/25.
DR P-PSDB: AAE00221.
XX
PT New variants of the gamma subunit of vertebrate adenosine
PT monophosphate-activated kinase for diagnosis or treatment of disorders
PT associated with energy metabolism such as diabetes, obesity, and
PT myopathy.
XX
PS Claim 12; Fig 2; 71pp; English.
XX
CC The present sequence is a cDNA encoding human adenosine monophosphate
CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
CC PRKAG3. Mutation in PrKag3 results in an altered regulation of
CC carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
CC useful as therapeutic for treating carbohydrate metabolism disorders such
CC as diabetes, obesity, and disorders associated with muscle metabolism
CC such as myopathy and cardiovascular diseases, to modulate AMPK
CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
CC and its functionally altered mutants are useful for the diagnostic
CC evaluation, genetic testing and prognosis of a metabolic disorder.
CC preferably a carbohydrate metabolism disorder. Primers that can detect
CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
CC useful for detecting a dysfunction of carbohydrate metabolism resulting
CC from the expression of a functionally altered allele of PRKAG3.
CC Transgenic animal and host cell transformed with PRKAG3 or a
CC heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC screening compounds able to modulate AMPK activity. Nucleic acid
CC encoding PRKAG3 is useful for detecting mutations in a PrKag3 gene, or
CC in a sequence encoding the first cystathione beta synthase (CBS) domain
CC of PRKAG3 and is useful in gene therapy.
XX
SQ Sequence 2109 BP; 458 A; 621 C; 560 G; 470 T; 0 other:

Query Match 87.98; Score 1447; DB 22; Length 2109;
Best Local Similarity 97.38; Pred. NO. 0;
Matches 1495; Conservative 0; Mismatches 35; Indels 7; Gaps 2;

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Oy 161 agaatcgttggaaaacggaaggccaaagccttgagatggaacaggaagtcggtggaag 220
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RESULT 4

AAD03321

ID AAD03321 standard; DNA; 2022 BP.

XX AC

AAD03321;

DT 13-JUN-2001 (first entry)

XX DE

Sus scrofa PRKAG3 splice variant DNA.

XX KW

Pig: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
 PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
 genetic testing; carbohydrate metabolism disorder; skeletal muscle;

cyathione beta synthase; CBS; cardiact; gene therapy; ds.

XX OS

Sus scrofa.

XX FH

Key Location/Qualifiers

FT CDS

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W0200120003-A2.

PD 22-MAR-2001.

PF 11-SEP-2000; 2000MO-EP09896.

PR 10-SEP-1999; 99EP-0402236.

PR 18-MAY-2000; 2000EP-0401388.

XX PA

(INRG) INRA INST NAT RECH AGRONOMIQUE.

PA (ANDE/) ANDERSSON L.

PA (LOOF/) LOOF C.

PA (KALM/) KALM E.

PI PI

Andersson L, Loof C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
 Iannuccelli N, Gellin J, Le Roy P, Chardon P;

DR DR

WPI: 2001-244810/25.

DR P-PSDB; AAEO0224.

XX PT

New variants of the gamma subunit of vertebrate adenosine
 monophosphate-activated kinase for diagnosis or treatment of disorders

PT PT

associated with energy metabolism such as diabetes, obesity, and
 myopathy -

XX PS

Claim 12; Page 69; 71pp; English.

XX CC

The present sequence is pig adenosine monophosphate (AMP)-activated
 kinase (AMPK) gamma subunit muscle-specific isoform, PRKAG3 splice

variant DNA. Prkag3 gene is located in the RN locus of chromosome 15.

Mutation in Prkag3 results in an altered regulation of carbohydrate

metabolism, particularly in skeletal muscle. PRKAG3 is useful as

therapeutic for treating carbohydrate metabolism disorders such as

diabetes, obesity, and disorders associated with muscle metabolism
 such as myopathy and cardiovascular diseases, to modulate AMPK

activity, and for restoring a normal AMPK function. PRKAG3 sequence
 and its functionally altered mutants are useful for the diagnostic

evaluation, genetic testing and prognosis of a metabolic disorder,

CC preferably a carbohydrate metabolism disorder. Primers that can detect
CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
CC useful for detecting a dysfunction of carbohydrate metabolism resulting
CC from the expression of a functionally altered allele of PRKAG3.
CC Transgenic animal and host cell transformed with PRKAG3 or a
CC heterodimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC screening compounds able to modulate AMPK activity. Nucleic acid
CC encoding PRKAG3 is useful for detecting mutations in a Pfkfb3 gene, or
CC in a sequence encoding the first cystathionine beta synthase (CBS) domain
CC of PRKAG3 and is useful in gene therapy.

Sequence 2022 BP; 412 A; 623 C; 593 G; 394 T; 0 other;

Query Match	71.28	Score	1172.8	DB	22	Length	2022
Best Local Similarity	85.78	Pred. No.	4e-296				
Matches 1351; Conservative	0	Mismatches	217	Indels	8	Gaps	4

[illegible]

QY	890	gaagacgcgtttgaagctggtctaaacccctcaagaagccggtatccatctgcctgctt	949
Db	946	gacacgcctgtcttcgaagctgtgtctaaagccctcatcaaaagacccggtatccacccgctgcgcgttc	1005
QY	950	cttgaccggcggtcaaggccaagatctccatatctctcaacacaaagccctgtccaattc	1009
Db	1006	ctggaccctgtctctccgggggctgtgtctccacaatcccaacaataagcggtcttccaagtcc	1065
QY	1010	ctggacaacttttggcttcaccgtctgcaccccgccctctctctctcaaccgacatcccaagat	1065
Db	1066	ctggacaacttttggcaccctcgtgcaccccgccctctctctctcaaccgacatcccaagat	1122
QY	1070	ttgggacatccgacacatcccgagacttgctgtgtgtctgtgagagacagaccacatctgact	1129
Db	1126	ttgggacatccgacacatcccgagacttgctgtgtgtctgtgagagacagaccacatctgact	1189
QY	1130	gcacgtggacacatctttgtggagaccggcggtgtgtctgcacatgcctgcgtgtctaaagatgtgt	1189
Db	1186	gcacgtggacacatcttcggtgagaccggcggtgtgtctgcacatgcctgcgtgtctaaagatgtgt	1245
QY	1190	cagcttcgtgggcccctctatctcccgctctgtatgtatctcaaccctgctgcacgacaacatcac	1245
Db	1246	cagcttcgtgggcccctctatctctgcgtctgtatgtatctcaaccctgctgcacgacaacatcac	1309
QY	1250	aaccacctgtgacatgtagtggtgtggagaaagccctgtaggtcagagagacatactgtctggagga	1309
Db	1306	aaccacctgtgacatgtagtggtgtggagaaagccctgtaggtcagagagacatactgtctggagga	1365
QY	1310	gtcccttctctgtccagcccccaaggagctcttggtggaaagtatccgacaggaatgtctccggag	1365
Db	1366	gtcccttctctgtccagcccccaaggagctcttggtggaaagtatctgtgacccggagatgtctccggag	1422
QY	1370	caggtgacacaggtgcgt	1422
Db	1426	caggtgtgacccgcctgt	1488
QY	1430	tccgacatctcttcagggcactgtgtgtctcagggccctgtctgtgacatgtgacctcggggctgta	1489
Db	1486	tctgacatctcttcagggcctctgt	1545
QY	1490	gaagctctgaagtcctctcaatcccaagaccact-gcacacctgtgaaagccaaatgaagggaact	1548
Db	1546	gaacctgtgaaaccttgcctctcaagaccgacacctgtgaaagccaaatgaagggaagc-	1604
QY	1549	ggagaaactcaagctctctctcccccacccccaattgtctgtgtcaagctatgtatcaagta	1608
Db	1605	cgtgtgacccagctctctcaatctccctcaagccccaactgtctgtgtctgtctgttctgtatgta	1666
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Db	1665	ggctctgtcccggggccc	1680

	Key	Location/Qualifiers
RESULT	5	
AAD03319		
ID	AAD03319	standard; cDNA; 1873 BP.
XX		
AC	AAD03319;	
XX		
DT	13-JUN-2001	(first entry)
XX		
DE	Pig	AMPK gamma subunit muscle-specific isoform, complete PRKAG3 cDNA.
XX		
KW	Pig;	gamma subunit; adenosine monophosphate-activated kinase; AMPK;
KW	PRKAG3;	diabetes; obesity; myopathy; cardiovascular disease; anorectic
KW	genetic testing;	carbohydrate metabolism disorder; skeletal muscle;
KW	cyathione beta synthase; CBS;	cardiant; gene therapy; RN locus;
KW	chromosome 15;	ss.
XX		
OS	Sus	scrofa.
XX		
FH		

Dh 1258 gaccgagatgtctcgaggaacaggtgacccgctgtgtctgtgatatgagaccagacact 1317
Oy 1412 ttggagctgtgtctccctctccgacatcttcacaggaactgtgtcagccctgtgtgcatc 1471
Dh 1318 ctggcgctgtgtgtctctgtacatcttcacaggaactgtgtcagccctgtgtgcatc 1377
Oy 1472 gatccctcgagggctctgagagaatctgtatctccatcccaagaccact-gaacactgtg 1530
Dh 1378 gatgctctcgagggctctgagagaatctgtatctccatcccaagaccactgtgacactgtg 1437
Oy 1531 aagcacaatgaaggaactgtgagaaactcagctcctcctcccaagcccatgtgtgtgt 1590
Dh 1438 aagcacaatgaaggaagc-cgtgtgactcagctcctcctcccaagcccatgtgtgtgt 1496
Oy 1591 tcaactatgactcaggtgagctcgtccctgtggcc 1624
Dh 1497 ctgtctctgtctcagtgagctcgcgggggccc 1530

RESULT 6

AA03295
ID AAD03295 standard; cDNA; 1867 bp.

AC AAD03295;

DT 13-JUN-2001 (first entry)

DE Pig AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.

XX
KW Pig; gamma subunit; adenosine monophosphate-activated kinase; AMPK;
KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
KW cystathione beta synthase; CBS; candidate; gene therapy; RN locus;
KM chromosome 15; ss.

XX
OS Sus scrofa.

XX
FH Key Location/Qualifiers

FT 5'UTR 1..471

FT CDS /*tag- a 472..1389

FT /*tag- b /product- "Sus scrofa PRKAG3 protein"
FT 3'UTR 1390..1867

FT /*tag- c

XX
PN MO200120003-A2.

XX
PD 22-MAR-2001.

XX
PF 11-SEP-2000; 2000MO-EP09896.

XX
PR 10-SEP-1999; 99EP-0402236.

XX
PR 18-MAY-2000; 2000EP-0401388.

XX
PA (INRG) INRA INST NAT RECH AGRONOMIQUE.

XX
PA (ANDE/) ANDERSSON L.

XX
PA (LOOF/) LOOF C.

XX
PA (KALM/) KALM E.

XX
PI Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;

XX
PI Iannuccelli N, Gellin J, Le Roy P, Chardon P;

XX
DR MPI: 2001-244810/25.

XX
DR P-PSDB; AAE00220.

XX
PT New variants of the gamma subunit of vertebrate adenosine
PT monophosphate-activated kinase for diagnosis or treatment of disorders
PT associated with energy metabolism such as diabetes, obesity, and
PT myopathy -
XX
PS Claim 12: Fig 2; 71pp; English.

CC The present sequence is a cDNA encoding pig adenosine monophosphate
CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
CC PRKAG3. Prkag3 gene is located in the RN locus of chromosome 15.
CC Mutation in Prkag3 results in an altered regulation of carbohydrate
CC metabolism, particularly in skeletal muscle. PRKAG3 is useful as
CC therapeutic for treating carbohydrate metabolism disorders such as
CC diabetes, obesity, and disorders associated with muscle metabolism
CC such as myopathy and cardiovascular diseases, to modulate AMPK
CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
CC and its functionally altered mutants are useful for the diagnostic
CC evaluation, genetic testing and prognosis of a metabolic disorder,
CC preferably a carbohydrate metabolism disorder. Primers that can detect
CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
CC useful for detecting a dysfunction of carbohydrate metabolism resulting
CC from the expression of a functionally altered allele of PRKAG3.
CC Transgenic animal and host cell transformed with PRKAG3 or a
CC heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC screening compounds able to modulate AMPK activity. Nucleic acid
CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
CC in a sequence encoding the first cystathione beta synthase (CBS) domain
CC of PRKAG3 and is useful in gene therapy.

XX
SQ Sequence 1867 bp; 380 A; 583 C; 529 G; 375 T; 0 other;

Query Match 68.9%; Score 1134.4; DB 22; Length 1867;
Best Local Similarity 85.7%; Pred. No. 4.1e-286;
Matches 1309; Conservative 0; Mismatches 211; Indels 8; Gaps 4;

Oy 101 ttcttagacaagaagaacagcagctatgtccatccagcgtgtgacagcagcattcaga 160
Dh 1 ttcttagacaagaagaagcggttcaatgtccatccagcgtgtgacacccagcagaa 60
Oy 161 agaatccgttgggaacaggaagcgaagccttctgagatgagacagaagcagatcggtgag 220
Dh 61 agaacatctgggaacaggaagcgaagccttctgagatgagacagaagcagatcggtgag 120
Oy 221 gaagggagagaccagcagtgtaagggaggtcccggtccagcagcagctgtatgtgtccac 280
Dh 121 gaagggagagaccagcagtgtaagggaggtcccggtccagcagcagctgtatgtgtgtccac 180
Oy 281 gggcttgagagcacaattccccaagacacaccccttggctcagctgtatgtgtgtgtgt 337
Dh 181 gggcagagagcacaattccccaagcagcagccttggctcagcagcagcgtgtgtgtgtgtgt 240
Oy 338 gtgggacatcccaacagaggtgtggactgtccctccctctgactgtacagcctcaagctga 397
Dh 241 gtggacacacccccaacagagcgggacatccctccctctgactgtgtcagcctcaagcctcc 300
Oy 398 ggtccagcagacagatgtgtgagctgtgacaggaattccccaagcagcagcgtgtgtgtgt 457
Dh 301 gactcacaacagacacatctgtgacatctgtgacatgtgtgtgtgtgtgtgtgtgtgtgtgt 360
Oy 458 tgtgagctagaagagcctgtctggaagagagcctgtgctgtgtgtgtgtgtgtgtgtgtgtgt 517
Dh 361 gatgagct---tggtgtgtgtggaagagagcagcagcagcgtgtgtgtgtgtgtgtgtgtgtgt 417
Oy 518 ttcccaagctgt 577
Dh 418 ttcccaagctgt 477
Oy 578 ttcatgcaagagcacaacccgtctagatgagtgaggaactcagctcagctcagctcagctcagctc 637
Dh 478 ttcatgcaagagcacaacccgtctagatgagtgaggaactcagctcagctcagctcagctcagctc 537
Oy 638 gacacacatgtgagatcagaagaagccttctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 697
Dh 538 gacacacatgtgagatcagaagaagccttctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 597
Oy 698 gccctctatgtgagcagaagaagcagccttctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 757
Dh 598 gaccttctgtgagcagaagaagcagccttctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 657

QY	758	atctctgtctgtatctgcctactacacggtctcccccctggtccagacatctatgaagatctgaacaa	817	
Db	658	atcttt	717	
QY	818	catcaagaatctgaagaccctggagagagatctaacctgcgaagacctctctcaagacctgtctctcc	877	
Db	718	catcaagaatctgaagaccctggagagagatctaacctgcgaagacctctctcaagacctgtctctcc	777	
QY	878	atctctcttaatagataagctgttttgaagctgtctctaacacctcaatcaagaaccggaatccat	937	
Db	778	atctctctcccaatgaacagcctgttctgaagctgtctctaacacctcaatcaagaaccggaatccac	837	
QY	938	cgcctctgctctctctctaaccccggtgtcaagccaaagctgaacatccatctctcaacacaaagc	997	
Db	838	cgcctctgctctctctctaaccccggtgtctcaagccaaagctgaacatccatctctcaacacaaagc	897	
QY	998	ctctctcaagctctctctgaacatcttctgttctccctctgtctcccgccctctctctctctaacgc	1057	
Db	898	ctctctcaagctctctctgaacatcttctgttctccctctgtctcccgccctctctctctctaacgc	957	
QY	1058	atcatctcaagaatttgggcatctcgcaaatctccgaagaccttggcctgtgtgtctgtgagaacgca	1117	
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QY	1118	cccatctctgctctgcaactgtgacacatctctgtgtgaagccggtgtgtctctgcaactctctgtgct	1177	
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QY	1178	aacgaatctgtgtcaggttcgttggcctctatctcccgctttgatgtatcaatcaatctgtctgc	1237	
Db	1078	aacgaatctgtgtcaggttcgttggcctctatctctcgctttgatgtatcaatcaatctgtctgc	1137	
QY	1238	cagcaaaaacctacaacacccctctgagacatgaatgtgtgagaaagccctctgaagcaagagacata	1297	
Db	1138	caacaaacatacaacacccctctgagacatgaatgtgtgagaaagccctctgaagcaagagacatctg	1197	
QY	1298	tgctctgagagagatctctctctctctgacccccaagagaccttgggtggaaagtgaatcgacag	1357	
Db	1198	tgctctgagagagagatctctctctctctgacccccaagagaccttgggtggaaagtgaatcgacg	1257	
QY	1358	atgtctctgagagagagatctctgagacagagctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt	1417	
Db	1258	atgtctctgagagagagatctctgagacagagctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt	1317	
QY	1418	gtgtgtctctctctctctctgacacatctctcaagagctgtgtgtgtgtgtgtgtgtgtgtgtgtgt	1477	
Db	1318	gtgtgtctctctctctctctgacacatctctcaagagctgtgtgtgtgtgtgtgtgtgtgtgtgtgt	1377	
QY	1478	ctctgagagcctctgagaagaatctctgaatctctcaatctccaaagccacct-gaacacctggagaagca	1536	
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QY	1537	atgagaagaaactctgagaagaactctcaacatctctctctcccccacacacacatctgtctgaact	1596	
Db	1438	atgagaagaaactctgagaagaactctcaacatctctctctcccccacacacacacatctgtctgaact	1496	
QY	1597	atgatactcaagatgaagctctgtccctggagcc	1624	
Db	1497	atgatactcaagatgaagctctgtccctggagcc	1524	
RESULT 7				
ID	ABA08485	standard: cdna: 547 bp.		
AC	ABA08485:			
XX	11-JAN-2002	(first entry)		
XX	Human AMP-activated protein kinase subunit homologue cdna, SEQ ID NO:261			
XX	Human: cytokine: cell proliferation: cell differentiation: growth factor:			
KW	haematopoiesis regulation: tissue growth: immunomodulator: activin:			

KM inhibitin, chemotaxis; chemokines; thrombolysis; oncogenesis;
 KM proliferation; metastasis; cancer; tumor; haematopoietic disorder;
 KM myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;
 KM chronic inflammatory condition; proliferative retinopathy;
 KM atherosclerosis; coronary heart disease; arterial ischaemia;
 KM bone disorder; osteoporosis; vascular growth disorder;
 KM tissue regeneration; wound healing; infection; immune disorder;
 KM cell culture; drug screening; gene therapy; antiinflammatory;
 KM antisthmatic; antiarthritic; haemostatic; antiatherosclerotic;
 KM cytosolic; osteopathic; vasotropic; cardiatic; vitricide; antibacterial;
 KM antiungal; vulninary; antulcer; ss.
 XX
 OS Homo sapiens.
 XX
 XX NO200157188-A2.
 PN
 XX 09-AUG-2001.
 PD
 XX
 PF 05-FEB-2001; 2001WO-US03800.
 PR 03-FEB-2000; 2000US-0496914.
 PR 27-APR-2000; 2000US-0560875.
 PA (HYSE-) HYSEQ INC.
 XX
 XX
 PI Tang YT, Liu C, Drmanac RT;
 DR MPI: 2001-457740/49.
 DR P-PSDB; ABBI1241.
 XX
 PT Human proteins and DNA encoding sequences useful for preventing,
 PT treating or ameliorating a medical condition in a mammalian subject
 PT e.g. arthritis and cancer -
 XX
 PS Claim 1; Page 429; 1963pp; English.
 XX
 XX Sequences ABBI0981-ABBI2330 represent 1350 novel human polypeptides, and
 CC sequences ABA084225-ABA09574 represent nucleic acids encoding them. The
 CC invention also relates to vectors and recombinant host cells comprising a
 CC nucleotide of the invention, methods of producing the novel polypeptides,
 CC antibodies against the polypeptides, methods of detecting the nucleotides
 CC or polypeptides in a sample, and methods of identifying compounds which
 CC bind to polypeptides of the invention. Although novel, many of the
 CC polypeptides of the invention have homology to known proteins, thereby
 CC giving an insight into their probable biological activities, and hence
 CC potential therapeutic applications. The polypeptides of the invention may
 CC have various activities, including cytokine, cell proliferation or cell
 CC differentiation activities; stem cell growth factor activity;
 CC haematopoietic regulatory activity; tissue growth factor activity;
 CC immunomodulatory activity; activin or inhibin-related activities;
 CC thrombolytic or chemokine activities; haemostatic, thrombotic or
 CC thrombolytic activities; receptor or ligand activities; or may be
 CC involved in oncogenesis, cancer cell proliferation or metastasis.
 CC Depending on their biological activities, polypeptides and nucleotides of
 CC the invention are useful for preventing, treating or ameliorating medical
 CC conditions, e.g., by protein or gene therapy. Such conditions include
 CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell
 CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),
 CC proliferative retinopathy, atherosclerosis, coronary heart disease,
 CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal
 CC vascular growth. Polypeptides involved with tissue regeneration and
 CC repair (or nucleic acids encoding them) may be used to promote wound
 CC healing (e.g., of burns, incisions and ulcers), while those with
 CC immunomodulatory activities may be used in the treatment of viral,
 CC bacterial and fungal infections in addition to immune disorders.
 CC Polypeptides with growth factor activity may be used in cell cultures to
 CC promote cell growth. For example, such polypeptides may be used to
 CC manipulate stem cells in culture to give rise to neuroepithelial cells
 CC that can be used to augment or replace cells damaged by illness,
 CC autoimmune disease or accidental damage. The polypeptides and nucleotides
 CC may also be used in the diagnosis of the above conditions, and in drug
 CC screening techniques. The present sequence represents a cDNA encoding a
 CC novel human polypeptide of the invention.

[illegible]

XX	Disease associated protein kinase DAPK-7 CDNA.
XX	
DE	DAPK-7; disease associated protein kinase; human; diagnosis;
KW	therapy; adult respiratory distress syndrome; allergy; asthma;
KM	arteriosclerosis; bronchitis; emphysema; hypereosinophilia;
KM	myocardial infarction; pericardial inflammation; anaemia;;
KM	rheumatoid arthritis; Addison's disease; AIDS; atherosclerosis;
KM	atopic dermatitis; dermatomycosis; diabetes mellitus;
KM	glomerulonephritis; gout; Grave's disease; lupus erythematosus;
KM	multiple sclerosis; myasthenia gravis; osteoarthritis;
KM	osteoporosis; pancreatitis; polycystic kidney disease;
KM	polymyositis; scleroderma; Sjogren's syndrome;
KM	autoimmune thyroiditis; cancer; infection; trauma;
XX	cell proliferation; ss.
OS	
XX	Homo sapiens.
FT	
CD	Key Location/Qualifiers
FT	265..1503
FT	/tag- a
XX	
FN	WO9858052-A2.
XX	
PD	23-DEC-1998.
XX	
PF	19-JUN-1998; 98WO-US12813.
XX	
PR	19-JUN-1997; 97US-0878989.
XX	
PA	(INCY-) INCYTE PHARM INC.
PI	Bandman O, Corley NC, Goli SK, Guegler KJ, Hillman JL;
PI	Lal P, Shah P;
XX	
WP	1999-080953/07

DR P-PSDB: AAM88438.
XX New disease associated protein kinases - used to stimulate cell
PT proliferation and to treat the immune response and cancer
XX
PS Claim 5: Page 66-67; 93pp; English.

XX This cDNA sequence codes for human disease associated protein
CC kinase DAPK-7 (see AAM88438). DAPK-7 cDNA was first identified in
CC the PENITUT01 cDNA library using a computer search for amino acid
CC alignments, and a consensus sequence was derived from the extended
CC and overlapping Incyte clones 3075712/HEARNOT01, 842220/PROST05,
CC 1364747/SCORONOT2, 145972 and 145802/PENITUT01 and 1479332/CORPOT02.
CC DAPK-7 shows 73% homology with the human foetal liver Apk gamma
CC subunit (GI 1335856), and is associated with cDNA libraries which
CC are immortalised or cancerous and show inflammatory or immune
CC responses. The invention provides disease associated protein kinases
CC DARK-1 to DARK-7 (see AAM88433-38) and cDNA clones encoding them (see
CC AAM06831-36 and AAM06882), as well as expression vectors, host cells,
CC agonists, antagonists and antibodies. The invention further provides
CC uses of such products in the diagnosis, prevention and treatment of
CC diseases associated with cell proliferation, especially cancer or an
CC immune response (claimed). Conditions that may be treated include
CC adult respiratory distress syndrome, allergies, asthma,
CC arteriosclerosis, bronchitis, emphysema, hypercosinophilia,
CC myocardial or pericardial inflammation, rheumatoid arthritis,
CC Addison's disease, AIDS, anaemia, atherosclerosis, various diseases
CC of the digestive system, atopic dermatitis, dermatomyositis, diabetes
CC mellitus, glomerulonephritis, gout, Grave's disease, lupus
CC erythematosus, multiple sclerosis, myasthenia gravis,
CC osteoarthritis, osteoporosis, pancreatitis, polycystic kidney
CC disease, polymyositis, scleroderma, Sjogren's syndrome, autoimmune
CC thyroiditis, complications of cancer, extracorporeal circulation,
CC viral, bacterial, fungal, parasitic, protozoal and helminthic
CC infections, and trauma (disclosed). The DAPK nucleic acids are
CC also used in a method for detection of DAPK expression levels in a
CC biological sample.
XX
XX

SQ Sequence 1435 BP; 421 A; 298 C; 331 G; 385 T; 0 other:

Query Match 22.3%; Score 366.6; DB 20; Length 1435;
Best Local Similarity 62.9%; Pred. No. 1.1e-85;
Matches 567; Conservative 0; Mismatches 334; Indels 0; Gaps 0;

OY 567 tctaatgctctcatgcaagagacacccctcgtcagatgcagcactgctccaagc 626
DB 183 tttatctgcatgcatgaggtacacacagtgltacatcgcttccaacagcttcaagc 242
OY 627 tagtcatctcgacacccctgctggagatcaagaagcctcttgcctgtgggccaagc 686
DB 243 ttgtgtcttcttgatctcatcaatgaagtaaaagcctcttgcctgtgggccaagc 302
OY 687 gttgtgggagcaccctctatggaagaagaagcagcttctgtgggagtcgcaaca 746
DB 303 ggttcgcagcagcagcagcagctgtggagagtaaaacaaagtttctaggaatgcaaca 362
OY 747 tcaatgacttcatctgtgtcgtacgtcactacaagttcccccgtgtccagatctatg 806
DB 363 ttacagattctctaaatctactacatagatactaaatcaccctatggtacagattatg 422
OY 807 agatgaacaacaataaattgagccttgaggagagatcactacctaagaagtcgctcact 866
DB 423 aattgaggaacaataaattggaacatgaggagagccttatttacaagaatacattaaagc 482
OY 867 ctctgttccatctctctactaatagtatgctgttgaagctgttcacacccctcatcaaga 926
DB 483 ctcttgatgaatatactctcagatgaagcctcttcgattgctgtatatacctcttgatcaaa 542
OY 927 accgcatccatgcgctgtcttgcagccgggtgtcaggaagaagtaactccatctcca 986
DB 543 ataaaatccacagatgtcccgcttaccctcatcagtggtggaatgacttatatactta 602

OY 987 cacacaagccttgcataagttctctgcacatcttggcttccctgtgcgccgcctcct 1046
DB 603 ccccaaaaagaatctctcaagttctctccagcgttcttatctgcatatgcacaaagcctct 662
OY 1047 tccctaccgacatctcaaatcttggcatcgcgcacatctccgacttgcgtgtggtgc 1106
DB 663 tcatgaagcagaacccttgatgagcttggaataggaacgtaaccacaacatctcctcaaac 722
OY 1107 tggagacagcaccatctgacttgcacttgcacatcttctgtgaacggcggtgtgtcac 1166
DB 723 atcagaacatcccatctcatcaagaagccttgaaactatcttggtaagaagcgaatatacagctc 782
OY 1167 tgcctgtgttcaagaatgtgtcgaagtcgtggcctctatctccgcttggatgtatc 1226
DB 783 tgcctgtgttggatgagtcaggaagaatgtgtgatatattatccaaatlttgatgaa 842
OY 1227 accggcgtgcccaacacccatccacacacccctggacatgagtggtgggaacccctggagc 1286
DB 843 atcttgcgtcgtgagaataacatacaataaactagatatacaggtgaacccagccttcaagc 902
OY 1287 agaggaacatagtctgtgaggagcttcttccgtccagcccaagagacttgggggaag 1346
DB 903 accgttcacagatatttgaagtggtgtggaagtgcataaagcttgaaatactgagacaa 962
OY 1347 tgcacgaagaatgtctcgtggagcaggtacacagcgtgtgtgtgtgagcagagcccaagc 1406
DB 963 tctgtgacaagaatgataagaagctgaggtccatcgctgtgtgtgtgtaataatgaagcagala 1022
OY 1407 atccttgggggtgtgtcctctccgcacatccctcagagcagctgtgtccagcctctgt 1466
DB 1023 gtattgtgggtatatttccctctcgcgacatcttcgcaagccctgatctccacacagcag 1082
OY 1467 g 1467
DB 1083 g 1083

RESULT 12
AAH14839
ID AAH14839 standard; cDNA: 2223 BP.
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AC AAH14839;
XX
DT 26-JUN-2001 (first entry)
XX
DE Human cDNA sequence SEQ ID NO:12860.
XX
KW Human; primer: detection; diagnosis; antisense therapy; gene therapy; ss.
XX
OS Homo sapiens.
XX
FN EP1074617-A2.
XX
PD 07-FEB-2001.
XX
PE 28-JUL-2000; 2000EP-0116126.
XX
PR 29-JUL-1999; 99JP-0248036.
XX
PR 27-AUG-1999; 99JP-0300253.
XX
PR 11-JAN-2000; 2000JP-0118776.
XX
PR 02-MAY-2000; 2000JP-0183767.
XX
PR 09-JUN-2000; 2000JP-0241899.
XX
XX (HELI-) HELIX RES INST.
XX
XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
XX WPL: 2001-318749/34.
XX
XX primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the

XX Sequence 3261 BP; 810 A; 952 C; 931 G; 568 T; 0 other;
50

Query Match	18.2%	Score 299.2;	DB 23;	Length 3261;
Best Local Similarity	58.6%	Pred. No. 6.1e-68;		
Matches 539; Conservative	0;	Mismatches 378;	Indels 3;	Gaps 1;

[illegible]

AAH43684	RESULT_14
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AC	AAH43684;
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DT	21-JAN-2002 (first entry)
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DE	PRKAG3 intron 10 - 3'UTR.
XX	
KW	Human: AMP-activated protein kinase gamma 3 subunit; PRKAG3; variant; metabolic disease; diabetes; obesity; substitution; ds.
XX	
OS	Homo sapiens.
XX	
FH	Key
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FT	857..1014
FT	/tag= g
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18-OCT-2001.	
XX	
06-APR-2001; 2001WO-SE00765.	
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07-APR-2000; 2000US-195665P.	
XX	
(AREX-) AREXS AB.	
PA	
XX	
PI	Andersson L, Luthman H, Marklund S;
XX	
DR	WPI: 2001-657170/75.
XX	
New variants of human AMP-activated protein kinase gamma3 subunit associated with a metabolic disease e.g. diabetes or obesity and method for determining a risk estimate of diseases In subject by detecting the variant -	
XX	
Example 1; Fig 4; 25pp; English.	
PS	
XX	
The sequences given in AAH43681-84 represents genomic fragments encoding the human AMP-activated protein kinase gamma 3 subunit (PRKAG3). Detecting the presence of the PRKAG3 DNA, or a variant, is useful in determining a risk estimate of a metabolic disease, such as diabetes or obesity, in a subject. The variation may occur in exons 3, 4 or 10. In exon 3 variation may be a substitution of a G for a C at nucleotide 320, resulting in the amino acid substitution P71A; in exon 4 variation may be a substitution of a T for a C at nucleotide 550; and in exon 10 variation may be a substitution of a T for a C at nucleotide 1037, resulting in the amino acid substitution R340W. There may also be nucleotide variation in Intron 6.	
Sequence 1014 BP; 192 A; 325 C; 271 G; 226 T; 0 other:	

Query Match 16.9%; Score 278.8; DB 22; Length 1014;
Best Local Similarity 97.6%; Pred. No. 8.3e-63;
Matches 283; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

Oy 1358 attgctcggaggaaggaaggaagctggtctgtctgtgagagagccagacatctcttgggc 1417
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Db 725 attcgagaccacaggaagcaagcttggctgtgtgagcagagccagacatctcttgggc 784
1418 gtgtgtctccctctccgacatctctcagcagctgtgtctcagccctgtgcatcgatgcc 1477
|||||
Db 785 gtgtgtccctctccgacatctctcagcagctgtgtctcagccctgtgcatcgatgcc 844
1478 ctgaggagcgcgaagaagatcctggaatctcaatcccaagccacctgcacacctggaagccaa 1537
|||||
Db 845 ctcgaggcctcgagaagatctggaatctcaatcccaagccacctgcacacctggaagccaa 904
1538 tgaagggaactggaagactcagcctcattctcccccaccccatctgtgtcagagcta 1597
|||||
Db 905 tgaagggaactggaagactcagcctcattctcccccaccccatctgtgtcagagcta 964
1598 tgaatcagggaagctctgcccctgtggccaagacacagcctctatgctctc 1647
|||||
Oy 965 tgaatcagggaagctctgcccctgtggccaatgacacagcctctatgctctc 1014
Db

RESULT 15
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ID AAK72740 standard; DNA: 3425 BP.

XX AAK72740:
XX
DT 06-NOV-2001 (first entry)
XX
DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:27552.
XX
KW Human; Immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX cytostatic; gene therapy; vaccine; metastasis; ds.
XX
OS Homo sapiens.
XX
PN MO200157182-A2.
PD
XX 09-AUG-2001.
PF
XX 17-JAN-2001; 2001MO-US01354.
XX
PR 31-JAN-2000; 2000US-0179065.
PR 04-FEB-2000; 2000US-0180628.
PR 24-FEB-2000; 2000US-0184664.
PR 02-MAR-2000; 2000US-0186350.
PR 16-MAR-2000; 2000US-0189874.
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PR 18-APR-2000; 2000US-0198123.
PR 19-MAY-2000; 2000US-0205515.
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PR 28-JUN-2000; 2000US-0214886.
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PR 26-JUL-2000; 2000US-0220963.
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PR 13-OCT-2000; 2000US-0239935.
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PR 01-NOV-2000; 2000US-0244617.
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QY	908	gtctcaaccctcatcaagaagccggatccatctgcctgcctgctctcttgagccgggtgtcagc	967
Db	302	gtctctttcatTTAAATTCGMAACAAGATCCACAGCCTGCCAGTTATTCACCAAAATCAGGC	361
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QY	1028	ctctgtcccccggccctcctctctctacgcacatccaaagatttggaagtcggacacalc	1087
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QY	1208	tccgcctcttgatgtgattcaccctgtgcctccagcaaacctacaaccctgtgacatgagct	1267
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QY	1328	caagcagagctgtggggaagltgacagagagaltg-ctcggagagcaggtgtacacagagctgtg	1386
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Db	781	actgtgtgattgaaatgattgtgtctcagagacatgt-TCACTGTCTGACATCTCTGACGCG	839
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ACCESSION	AL519198		
VERSION	AL519198.1	GI:12782691	
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SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.		
TITLE	Li, W.B., Gruber, C., Jessup, J. and Polayes, D.		
JOURNAL	Full-length cDNA libraries and normalization		
COMMENT	Unpublished (2001)		
FEATURES	Genoscope - Centre National de Sequencage		
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vector. Library is not normalized, but is the control for
the normalized libraries. Library was constructed by Life
Technologies. Contact : Feng Liang Life Technologies, a
division of Invitrogen 9800 Medical Center Drive Rockville
, Maryland 20850, USA Fax : (1) 301 610 8371 Email :
fliang@lifetech.com URL :
http://fulllength.invitrogen.com"
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Query Match      20.8% Score 343: DB 9: Length 826:
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OY 627 tagtcatctgcacacatgcagatgcacacacacacacacacacacacacacac 686
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DB 128 TGTGTATTTGATGATGACGCTCCCTGACGCTGACGCTGATGCTGATGCTGAT 187

OY 687 gtagcgagcagacacacacacacacacacacacacacacacacacacacacac 746
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OY 747 tcaactgactcaccacacacacacacacacacacacacacacacacacacacac 806
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DB 248 TCACTGATTTTCATCAATATCTGTCACCGCTACATTAATCACTTGTGATCACTTA 307

OY 807 agatlgacaacataagatltgagacacacacacacacacacacacacacacacac 866
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DB 308 ACCTAGACAGACACACAGATGAGAACTGGAGAGAGGTGATCTCCAGCACTCTTTA 367

OY 867 ctctgctccatctctctctctctctctctctctctctctctctctctctctca 926
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OY 1107 tggagacacacacacacacacacacacacacacacacacacacacacacacac 1166
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OY 1167 tgcctgtgtcaacagatgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 1226
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DB 668 TCCAGATGTGTGATGAGAAAGGCGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 727

OY 1227 accgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 1286
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AL513784
VERSION
AL513784.1 GI:12777278
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EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
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Li, W.B., Gruber, C., Jesssee, J. and Polayes, D.
Full-length cDNA libraries and normalization
Unpublished (2001)
COMMENT
Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 Evry cedex - France
Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.
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vector. Library was normalized. Library was constructed by
Life Technologies. Contact : Feng Liang Life Technologies,
a division of Invitrogen 9800 Medical Center Drive
Rockville, Maryland 20850, USA Fax : (1) 301 610 8371
Email : fliang@lifetech.com URL :
http://fulllength.invitrogen.com"
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OY 627 tagtcatctgcacacacacacacacacacacacacacacacacacacacacac 686
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OY 687 gtagcgagcagacacacacacacacacacacacacacacacacacacacacac 746
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OY 747 tcaactgactcaccacacacacacacacacacacacacacacacacacacacacac 806
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OY 807 agatlgacaacataagatltgagacacacacacacacacacacacacacacacacacac 866
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OY 867 ctctgtgtcctatctctctctctctctctctctctctctctctctctctctctca 926
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 401 CTTTATGATATATATCTCCAGATCCAAAGCCTTCTGATGCTGTATCTCTTGTAT 460

OY 927 accgagatcagcctgctgtctgtacacacacacacacacacacacacacacacac 986
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
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Query Match	23.7%	Score 390.2	DB 3	Length 1576
Best Local Similarity	64.9%	Pred. No. 1.8e-93		
Matches 578; Conservative	0	Mismatches 313	Indels	0
Gaps				

Db	176	TGATATCTTCCTTACATGAAGTCATCCGTGCTATGACCTGATTCCACAAGCTCCAAAT	2351
Qy	627	taagatcctcgacacacatgctggaatcaagaagccctctctgctcgtggaacg	686
Db	236	TGGTTTATTATTATACGCTCCCTGACGAGGTGAAGAAAGCTTTTTCGTTGGTAGCTAACG	295
Qy	687	glttgaggagccgcctctctgagaccaagaagcagagctcttgaggatgctgacca	746
Db	296	CTCTACGACTGCCCTTTATGTGGATACATAAGAACAAAGTTTGTGGGATCCTGACCA	355
Qy	747	tcaagactcaatccctgctgctgacatcgactcaagatccccccctgctcagatcatg	806
Db	356	TCTCATTTTTCATCATATTCCTGCACCCCTACTATAATACAGCCTTGTTACATCTATG	415
Qy	807	agaatbaaacaalaagaatlgaagccctggaaggagatctactcgtgaagcttgctcaagc	866
Db	416	AGCTAGACAAACACAAAGTACGAACCTTGACAGAGGTCATTCACAGCATCCTTTAAAC	475
Qy	867	cctcgtgctcaatctctcctaataatgaagccgttatgagctgtcactaacctcataaga	926
Db	476	CCCTTGCTGCATTTCTCCTTAATGCACAGCTGTTTGATGCTGTCTCTCATTTAATTGCA	535
Qy	927	accgagatcatcgccgtcgtctcttgaccccggtgtcaaggcaagctacatccatccata	986
Db	536	ACAATATCCACAGCTGCGCACTTATTACCCAGATACAGCAATTTGTTGATCTCTCA	595
Qy	987	caacaaagccgtcgtcaagctccctgacatctcttgctccctcgtcccgccgacctct	1047
Db	596	CCCAACACCGCATTTCTGAAGTTCTCTCAAAATGTTTATACACTAGTTTCCCCAACCAAGCT	655
Qy	1047	tccctctaacgacatcacagaatttggatcttggaatctggacatccgaagcttgctgtgtgc	1106
Db	656	TCATTTCCAAAGTCTCTGGAMAGCTACAGATTGGACATATGCTATGATGGTTTC	715
Qy	1107	tggagaacgaacccatccctgaactgaactgaacatctcttgtagccgggtgtgtctgac	1166
Db	716	GCATACACACCCCGCTGTATGTGGCTCTGGGATTTTGTAGACGATCGAGCTCAAGCC	775
Qy	1167	tgcctgtgtcaagaatctggtgtcaagtggtgggtgggtctatcccgcttgatgtgatic	1226
Db	776	TGCCAGTGTGTGATGACAAAGGCCGTGTGGTGCATCTACGCAAGTTTGAATGTTATCA	835
Qy	1227	accgtgctgccagacaacctacaacaccctgtgacatgtagtctggagaagaagccctgagcg	1286
Db	836	ATTGTGACGACGAAAMAACCTTACAAACCTAGATGTATGTGACTAAAGCTTTGCAC	895
Qy	1287	agaagaacatactgtctggaggagatcccttcctgcgaagcccaagagagcttgagggaag	1346
Db	896	ATGCATCACATTACTTTGAGGGGTGTTCTCAAGCTCACTCGATGATAGCACTGTGAGACCA	955
Qy	1347	tgaatcagaagatgtctcgggaagcagtaacagagctgtgtctagtggaagagaccagc	1406
Db	956	TGATCAACACAGCTAGTGAAGCAGACGCTTACACCGACTTGTAGTGTGATMAAATATGC	1015
Qy	1407	atcctctgtgcgtgtgctccctctccgaaatccctcgaagctgtgacta	1457
Db	1016	TGGTCAAGGGAATTGTATACACTGTCTGACATCTCTGACATCTCTGACGCCCTGGTGTCTA	1066

RESULT 2
 US-08-878-989-14
 Sequence 14, Application US/08878989
 Patent No. 5885803
 GENERAL INFORMATION:
 APPLICANT: Bandman, Olga
 APPLICANT: Hillman, Jennifer L.
 APPLICANT: Corley, Neil C.
 APPLICANT: Guegler, Karl G.
 APPLICANT: Lal, Preeti
 APPLICANT: Goli, Surya K.
 APPLICANT: Shah, Purvi
 TITLE OF INVENTION: DISEASE ASSOCIATED PROTEIN

```

1  TITLE OF INVENTION:  KINASES
2  NUMBER OF SEQUENCES:  21
3  CORRESPONDENCE ADDRESS:
4  ADDRESSEE:  Incyte Pharmaceuticals, Inc.
5  STREET:  3174 Porter Drive
6  CITY:  Palo Alto
7  STATE:  CA
8  COUNTRY:  USA
9  ZIP:  94304
10 COMPUTER READABLE FORM:
11 MEDIUM TYPE:  Diskette
12 COMPUTER:  IBM Compatible
13 OPERATING SYSTEM:  DOS
14 SOFTWARE:  FASTSEQ for Windows Version 2.0
15 CURRENT APPLICATION DATA:
16 APPLICATION NUMBER:  US/08/878,989
17 FILING DATE:
18 CLASSIFICATION:  435
19 PRIOR APPLICATION DATA:
20 APPLICATION NUMBER:
21 FILING DATE:
22 ATTORNEY/AGENT INFORMATION:
23 NAME:  Billings, Lucy J J
24 REGISTRATION NUMBER:  36,749
25 REFERENCE/DOCKET NUMBER:  PF-0321 US
26 TELECOMMUNICATION INFORMATION:
27 TELEPHONE:  415-855-0555
28 TELEFAX:  415-845-4166
29 TELEX:
30 INFORMATION FOR SEQ ID NO:  14:
31 SEQUENCE CHARACTERISTICS:
32 LENGTH:  1435 base pairs
33 TYPE:  nucleic acid
34 STRANDEDNESS:  single
35 TOPOLOGY:  linear
36 IMMEDIATE SOURCE:
37 LIBRARY:  PEN1UT01
38 CLONE:  1452972
39 US-08-878-989-14

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US-08-878-989-14

Query Match	Score	DB 2;	Length
22.38;	366.6;		1435;

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Matches 567; Conservative 0; Mismatches 334; Indels 0; Gaps 0;

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OY	567	lctlaaagcgcttcaatgacagagacaacactgtcactagatgccaatagctccaagc	62
Db	183	TTTAAATCGGATTTATATAGTGTCACAAAGTGTTATGACATGTTCCAAACAGTTCAAAAC	24
OY	627	tagtcaatctgcacacacagctggagatcaagaagcctcttgcctcgggtggccaagc	68
Db	243	TTGTTGCTTTGATACACTTTCACAAGTTAAAAAGGCGCTTTTGGCTTTGGTAGGCCAACG	30
OY	687	gtctgcgagcagccctctatggacacgaagaacagagctgttgaggatgctgacga	74
Db	303	GTCGCCAGACAGCGCCACTGTGGAGAGTAAAAAACAAGTTTGTTAGCAATGCTTAACAA	36
OY	747	tcaactgacttcaatcccggtgcgtcgaatcgtcactacaagtcctccctcgtgccagatctatg	80
Db	363	TTTACAGATTTTCATAAATAATACACTACATGATCTACTTAAATCACCCTATGTATACAGATTTATG	42
OY	807	agacttgacaacataagattgagacctggaggagatctctccgcgaagcgtcctccaagc	86
Db	423	AATTAGAGGAACCTTAAATATTGGAACATGAGGAGGCTTTATTTACAAAGAAACATTTAAAGC	48
OY	867	ctctggtctccalcctccctaataatgataagcgtgttgaagctgtctacaccctccaaga	92
Db	483	CTTTAGGATATATATCTCCAGATCAAGCCCTTCGATGTGTATATCTTTGATCAAAAA	54
OY	927	accggaatcgcgcctgcgcgttcttgaacccggtgtccaggacaagtactccaacatccca	98
Db	543	ATTAATAATCCACAGATTTGGCCGTTATTTACACCTTACAGTGGGAATCCACTTTATTTACTTA	60

Accession	Sequence	Position
Oy	caacaaagcgcgtccaaagllctcgcgaacttggtccctcgtgcccgccctc	1046
Dh	CCCCAAAAGATCTTCAAGTCTCCAGCTTTTATATGTGTATATGCCAAAAGCGTCT	662
Oy	1047	1106
Dh	663	722
Oy	1107	1166
Dh	723	782
Oy	1167	1226
Dh	783	842
Oy	1227	1286

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:      REGISTRATION NUMBER: 36,749
:      REFERENCE/DOCKET NUMBER: PF-0321 US
:      TELECOMMUNICATION INFORMATION:
:      TELEPHONE: 415-855-0555
:      TELEFAX: 415-845-4166
:      TELEX:
:      INFORMATION FOR SEQ ID NO: 14:
:      SEQUENCE CHARACTERISTICS:
:      LENGTH: 1435 base pairs
:      TYPE: nucleic acid
:      STRANDEDNESS: single
:      TOPOLOGY: linear
:      IMMEDIATE SOURCE:
:      LIBRARY: PENITUT01
:      CLONE: 1452972
:      US-09-272-796-14

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RESULT 3
 US-09-272-796-14
 : Sequence 14, Application US/092272796
 : Patent No. 6207148
 : GENERAL INFORMATION:
 : APPLICANT: Bandman, Olga
 : APPLICANT: Hillman, Jennifer L.
 : APPLICANT: Corley, Neil C.
 : APPLICANT: Guegler, Karl G.
 : APPLICANT: Lal, Preeti
 : APPLICANT: Goli, Surva K.
 : APPLICANT: Shah, Purvi
 : TITLE OF INVENTION: DISEASE ASSOCIATED PROTEIN
 : TITLE OF INVENTION: KINASES
 : NUMBER OF SEQUENCES: 21
 : CORRESPONDENCE ADDRESS:
 : ADDRESSEE: Incyte Pharmaceuticals, Inc.
 : STREET: 3174 Porter Drive
 : CITY: Palo Alto
 : STATE: CA
 : COUNTRY: USA
 : ZIP: 94304
 : COMPUTER READABLE FORM:
 : MEDIUM TYPE: Diskette
 : COMPUTER: IBM Compatible
 : OPERATING SYSTEM: DOS
 : SOFTWARE: FASTSEQ for Windows Version 2.0
 : CURRENT APPLICATION NUMBER:
 : APPLICATION NUMBER: US/09/272,796
 : FILING DATE:
 : CLASSIFICATION:
 : PRIOR APPLICATION DATA:
 : APPLICATION NUMBER: 08/878,989
 : FILING DATE:
 : ATTORNEY/AGENT INFORMATION:
 : NAME: Billings, Lucy J J

QY	1347	lgalcgaaggaattgctcgaggagcagctacacagggcttggctgtaattgagacagagaccgcgc	1406
DB	963	TCGTGGACACATAGCTAAGACGCTGAGCTGCATCGCTCGTGTGCTAATGTAAAGACAGATA	10222
QY	1407	atcctctggagcgctgctctccctctctcgacacatcctctcaagcacttggctgcagccctctgc	1466
DB	1023	GTAATTGGGAGTAATATTTCCCTCTCGGACATATCTGTGAACCCCTGATCTCTCAOACGACGACG	10822
QY	1467	g 1467	
DB	1083	G 1083	

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1      RESULT 4
2      : Sequence 14, Application US/08232463
3      : Patent No. 5670367
4      : GENERAL INFORMATION:
5      : APPLICANT: DORNER, F.
6      : APPLICANT: SCHEIFLINGER, F.
7      : APPLICANT: FALKNER, F. G.
8      : TITLE OF INVENTION: RECOMBINANT FOULPOX VIRUS
9      : NUMBER OF SEQUENCES: 52
10     : CORRESPONDENCE ADDRESS:
11     : ADDRESSEE: Foley & Lardner
12     : STREET: 1800 Diagonal Road, Suite 500
13     : CITY: Alexandria
14     : STATE: VA
15     : COUNTRY: USA
16     : ZIP: 22313-0299
17     : COMPUTER READABLE FORM:
18     : MEDIUM TYPE: Floppy disk
19     : COMPUTER: IBM PC compatible
20     : OPERATING SYSTEM: PC-DOS/MS-DOS
21     : SOFTWARE: Patentin Release #1.0, Version #1.25
22     : CURRENT APPLICATION DATA:
23     : APPLICATION NUMBER: US/08/232.463
24     : FILING DATE:
25     : CLASSIFICATION: 435
26     : PRIOR APPLICATION DATA:
27     : APPLICATION NUMBER: US/07/935.313
28     : FILING DATE:
29     : APPLICATION NUMBER: EP 91 114 300.6
30     : FILING DATE: 26-AUG-1991
31     : ATTORNEY/AGENT INFORMATION:
32     : NAME: BENT, Stephen A.
33     : REGISTRATION NUMBER: 29,768
34     : REFERENCE/DOCKET NUMBER: 30472/114 IMMU
35     : TELECOMMUNICATION INFORMATION:
36     : TELEPHONE: (703)836-9300
37     : TELEFAX: (703)683-4109
38     : TELEX: 899149
39     : INFORMATION FOR SEQ ID NO: 14:
40     : SEQUENCE CHARACTERISTICS:
41     : LENGTH: 7218 base pairs
42     : TYPE: nucleic acid
43     : STRANDEDNESS: single
44     : TOPOLOGY: linear
45     : IMMEDIATE SOURCE:
46     : CLONE: pTZ9pt-F15
47     : US-08-232-463-14

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Query Match	3.08	Score 49	DB 1	Length 7218
Best Local Similarity	2.68	Pred. NO. 0.0021		
Matches 10	Conservative 220	Mismatches 155	Indels 0	Gaps 0

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0Y      864 agcctctgagctccatcctccctcaagatagcctgtttgaagctgtctacaccctcatca 923
      ||| | | : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db     1057 AGCTTCGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1116
0Y      924 agaacgagatcatalgcctgcctgcctctcttgaccgcggtgttcaggcaacgacttccatcc 983

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[illegible]

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RESULT 5
US-09-056-105-8
; Sequence 8, Application US/09056105
; Patent No. 6287569
; GENERAL INFORMATION:
; APPLICANT: KIPPS, THOMAS J.
; APPLICANT: WU, YUNCI
; TITLE OF INVENTION: VACCINES WITH ENHANCED INTRACELLULAR
; TITLE OF INVENTION: PROCESSING
; FILE REFERENCE: 233/221
; CURRENT APPLICATION NUMBER: US/09/056-105
; CURRENT FILING DATE: 1998-04-06
; EARLIER APPLICATION NUMBER: 60/043,467
; EARLIER FILING DATE: 1997-04-10
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 1022
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-056-105-8

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Query Match	2.58;	Score 41.2;	DB 4;	Length 1022;
Best Local Similarity	44.78;	Pred. No. 0.12;		
Matches 160;	Conservative 0;	Mismatches 198;	Indels 0;	Gaps 0;

OY 373 ctctcagcagcagcagcctcaagctctgcagaggtctccagccagatgatcttgaggtctgcagcag 43
 Db 299 caattgctgcagagcgaaccatgaggtgtccagcagccaaagaagagagagagccagac 35
 OY 433 gtctcccaagcccaagagctgcaggtgagtgtagctagaagagctctctggaagagagcctgc 49
 Db 359 ctgcgctcgaagcagagagctctctgtctccgaggaagcactcagtaacaagtgagatgttggc 41
 OY 493 cctgtgctgtctcccgacagcccaattcccaagctgggtctgagtgatgacgaacgcggaa 55
 Db 419 tcaatttctgtctcgcaagtactgagccaaagagcttctgtccaaagaagcagaaagctgg 47
 OY 553 accgcgcgcagcagctctcactgcagctctcatgtcaggaagacacctgcctacagctgcagc 61
 Db 479 gagagctacccaataattacaagcgtgtcttctcctgtgactcttcggcaaaagcctcagtc 53
 OY 613 aactagctccaagtgatgactcttcgcagacacatgtctgagataccaagaagcctctcttcg 67
 Db 539 cctgaagatgactcttctgcattgcagtcgaggaagatggacccgcgcagcaacactcaac 58
 OY 673 tctggtgcagcaagtgctgcggcagccctctatgtgagacgaagaagacgctt 73


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1 ZIP: 10022
2
3 COMPUTER READABLE FORM:
4
5 MEDIA TYPE: Diskette, 5.25 inch, 360 kb storage
6
7 COMPUTER: IBM
8
9 OPERATING SYSTEM: PC-DOS
10
11 SOFTWARE: Wordperfect
12
13 CURRENT APPLICATION DATA:
14
15 APPLICATION NUMBER: US/08/299,849B
16
17 FILING DATE: 1-SEPTEMBER-1994
18
19 CLASSIFICATION: 435
20
21 PRIOR APPLICATION DATA:
22
23 APPLICATION NUMBER: 08/037,230
24
25 FILING DATE: 26-MARCH-1993
26
27 PRIOR APPLICATION DATA:
28
29 APPLICATION NUMBER: PCT/US92/04354
30
31 FILING DATE: 22-MAY-1992
32
33 PRIOR APPLICATION DATA:
34
35 APPLICATION NUMBER: 07/807,043
36
37 FILING DATE: 12-DECEMBER-1991
38
39 PRIOR APPLICATION DATA:
40
41 APPLICATION NUMBER: 07/764,364
42
43 FILING DATE: 23-SEPTEMBER-1991
44
45 PRIOR APPLICATION DATA:
46
47 APPLICATION NUMBER: 07/728,838
48
49 APPLICATION NUMBER: 9-JULY-1991
50
51 PRIOR APPLICATION DATA:
52
53 APPLICATION NUMBER: 07/705,702
54
55 FILING DATE: 23-MAY-1991
56
57 ATTORNEY/AGENT INFORMATION:
58
59 NAME: Hanson, No. 5612201man D.
60
61 REGISTRATION NUMBER: 30,946
62
63 REFERENCE/DOCKET NUMBER: LUD 5355
64
65 TELECOMMUNICATION INFORMATION:
66
67 TELEPHONE: (212) 688-9200
68
69 TELEFAX: (212) 838-3884
70
71 INFORMATION FOR SEQ ID NO: 14:
72
73 SEQUENCE CHARACTERISTICS:
74
75 LENGTH: 2531 base pairs
76
77 TYPE: nucleic acid
78
79 STRANDEDNESS: single
80
81 TOPOLOGY: linear
82
83 MOLECULE TYPE: genomic DNA
84
85 FEATURE:
86
87 NAME/KEY: MAGC-41 gene
88
89 US-08-299-849B-14

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	Query Match	2.4%:	Score 39.6:	DB 1:	Length 2531:
	Best Local Similarity	44.4%:	Pred. No. 0.43:		
	Matches 159:	Conservative	0:	Mismatches 199:	Indels 0;
				Gaps	0:
OY	373 ctcctacgtctcagcgctcgaagtgcgagccctccagcacagatgatgttggactgtgccaaggaa	432			
Db	858 CACTTGTCTGGAGCGCAACCCAAATGAGCGTTTCAGCAGCCAAAGAAGAGAGGGGCCAAGCAC	917			
OY	433 gtcccagaaccagaagcgcttgtagtgtgagcttagaagcgctcgcgtggaagaagcgctgc	492			
Db	918 CTCGCCTACCCAGAGTGCTCTTGTCCGAGAACCTCAGTAACAAGTGCATGAGCTGGCG	977			
OY	493 ccctgtccctgtccccgcagagcccaattcccaagcttggtgtgataagaaactgcgaa	552			
Db	978 TCAATTCTCGTCCGCCAAGTATTCAGAGCCAAAGAGCTGGTCACAAAGCGAANAATCTGGA	1037			
OY	553 acccgcgcccaagatctacatgcgtlcatygaagagcacacctgtctaagatgcatalgpc	612			
Db	1038 GAGAGTCATCAAAAATTCAGACGCCGTGCTTTCGTGATCTTTCGGCAAAAGCTCCGAGTC	1097			
OY	613 aaactagctcccaagctatgtcatcttcgacacaatgtcgtagaatcaagaagcgcttccttgc	672			
Db	1098 CCTGAAAGATGATCTTTGGCATTTGACGTTGAAGGAAGTAGGACCCACACCACCAACTACAC	1157			
OY	673 tcctgttgccaacggtgtgtggtgcagagcccccctatctgtgacagcaagaaagcagagcttc	730			

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Db      1158  CcTtGcACcTgcTgcGcGcTTCcTATcATgcTgcCCTgcGTAATATcATcTCTT  1213

RESULT      9
US-08-142-368A-14
: Sequence 14, Application US/08142368A
: Patent No. 5925729
:
GENERAL INFORMATION:
:
APPLICANT: Boon-Falleur, Thierry; Van der Bruggen, Thierry;
APPLICANT: Van den Eynde, Beno t; Van Pel, Aline; De Plaeen, Etienne
APPLICANT: Lurquin, Christophe; Chomez, Patrick; Traversari, Celia
TITLE OF INVENTION: Tumor Rejection Antigen Precursors, Tumor
TITLE OF SEQUENCE: Rejection Antigens and Uses Thereof
NUMBER OF SEQUENCES: 26
:
CORRESPONDENCE ADDRESS:
:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
:
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
:
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/142,368A
FILING DATE: 02-MAY-1994
CLASSIFICATION: 435
:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/04354
FILING DATE: 22-MAY-1992
:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/807,043
FILING DATE: 12-DECEMBER-1991
:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/764,364
FILING DATE: 23-SEPTEMBER-1991
:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
APPLICATION NUMBER: 9-JULY-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/705,702
FILING DATE: 23-May-1991
:
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 5925729man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5253.4-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
:
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 2531 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
:
MOLECULE TYPE: genomic DNA
FEATURE:
NAME/KEY: MAGE-41 gene
US-08-142-368A-14

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Query Match	2.48	Score 39.6	DB 2	Length 2531
Best Local Similarity	44.48	Pred. NO. 0.43		
Matches 159	Conservative	0	Mismatches 199	Indels 0
			Gaps	0
QY	373	ctctgactgtacagcctctaagctgcagagctccagacagatggtggagcttggccaaga	432	
Db	858	CACGTGCTGGAGGCAACCCATGAGGCTTCCAGCAGCCAAAGAGGGGGCCAAAGAC	917	
QY	433	gttcccgagccagagagccttggaagtgtgaactgaagaagctctctgtgaagaagaagcttc	492	

Db 918 CTCGCTGACGAGATCCTTGTCCGAGAACACTCAGTACAGTACGATGACTTGGC 977
Qy 493 cctgtgcctgtcccccgcagcccaatcccaagctggcgtggaatgaagaaatgcgaa 552
Db 978 TCATTTTCTGCTCCGCAAGTATGAGCAGACGCTGGTTCACAAAGCAGAAATGCTGGA 1037
Qy 553 acccgagcccgacatcagctcagctcagcagcagcagcagcagcagcagcagcagc 612
Db 1038 GAGAGTCATCAAAATTTACAGCCGCTGCTTCTGATCTTTCGCAAGCCTCCGAGTC 1097
Qy 613 aactagctccaaagctagctcagcagcagcagcagcagcagcagcagcagcagcagc 672
Db 1098 CCTGAAGTATGATCTTGGCATTGACGTAGAGAGAGTGCACCCACACACACTTACAC 1157
Qy 673 tctgtgtgcacaaagctgtgcgagcagccctcctatgtggaagcagaagcagcagc 730
Db 1158 CTTGTTCACCTGCTGGGCTTTCTTATGATGAGCCTGCTGGTATATATCAGATCTTT 1215

RESULT 10
US-08-967-727-14
Sequence 14, Application US/08967727
Patent No. 6023474
GENERAL INFORMATION:
APPLICANT: Gaugler, B atrice; Van den Eynde, Beno t;
APPLICANT: van der Bruggen, Pierre; Boon-Fallour, Thierry
TITLE OF INVENTION: Isolated Nucleic Acid Molecules Coding For
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESS: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,727
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/037,230
FILING DATE: 26-MARCH-1993
APPLICATION NUMBER: PCT/US92/04354
FILING DATE: 22-MAY-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/807,043
FILING DATE: 12-DECEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/764,365
FILING DATE: 23-SEPTEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
FILING DATE: 9-JULY-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/705,702
FILING DATE: 23-MAY-1991
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 6023474man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5353
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 2531 base pairs
TYPE: nucleic acid
STRANDEDNESS: single

TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
FEATURE:
NAME/KEY: MAGE-41 gene
US-08-967-727-14

Query Match 2.4%; Score 39.6; DB 3; Length 2531;
Best Local Similarity 44.4%; Pred. No. 0.43;
Matches 159; Conservative 0; Mismatches 199; Indels 0; Gaps 0;

Qy 373 cctgtgcctgtcccccgcagcccaatcccaagctggcgtggaatgaagaaatgcgaa 432
Db 858 CACTTGTGAGGAGCAACCCATGAGGCTTCCAGCAGCCACAGAAAGAGGAGGCGCAACAC 917
Qy 433 gtcccgagcccgagcagcagcagcagcagcagcagcagcagcagcagcagcagc 492
Db 918 CTCGCTGACGAGATCTTGTTCGAGAGACCTCAGTAACAGCTGATGATGCTG 977
Qy 493 cctgtgcctgtcccccgcagcccaatcccaagctggcgtggaatgaagaaatgcgaa 552
Db 978 TCATTTTCTGCTCCGCAAGTATGAGCAGACGCTGGTTCACAAAGCAGAAATGCTGGA 1037
Qy 553 acccgagcccgacatcagcagcagcagcagcagcagcagcagcagcagcagcagc 612
Db 1038 GAGAGTCATCAAAATTTACAGCCGCTGCTTCTGATGATGATGATGATGATGATG 1097
Qy 613 aactagctccaaagctagctcagcagcagcagcagcagcagcagcagcagcagcagc 672
Db 1098 CCTGAAGTATGATCTTGGCATTGACGTAGAGAGTGCACCCACACACACTTACAC 1157
Qy 673 tctgtgtgcacaaagctgtgcgagcagccctcctatgtggaagcagaagcagcagc 730
Db 1158 CTTGTTCACCTGCTGGGCTTTCTTATGATGAGCCTGCTGGTATATATCAGATCTTT 1215

RESULT 11
US-08-037-230D-14
Sequence 14, Application US/08037230D
Patent No. 6235525
GENERAL INFORMATION:
APPLICANT: Gaugler, B atrice; Van den Eynde, Beno t;
APPLICANT: van der Bruggen, Pierre; Boon-Fallour, Thierry
TITLE OF INVENTION: Isolated Nucleic Acid Molecules Coding For
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESS: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/037,230D
FILING DATE: 26-MARCH-1993
APPLICATION NUMBER: 05/08/037,230D
FILING DATE: 26-MARCH-1993
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/04354
FILING DATE: 22-MAY-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/807,043
FILING DATE: 12-DECEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/764,364
FILING DATE: 23-SEPTEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
FILING DATE: 9-JULY-1991


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QY 1014 acatcttggtccctgctgcccgcctctctccaccgacatccaagatttg 1073
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Db 126 NNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNSY 67
QY 1074 gcatcgagac 1083
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Db 66 NNNNSNNNNY 57

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RESULT 14
US-09-244-796-17/C
: Sequence 17, Application US/09244796
: Patent No. 6281344
: GENERAL INFORMATION:
: APPLICANT: Szostak, Jack W.
: APPLICANT: Roberts, Richard W.
: TITLE OF INVENTION: SELECTION OF PROTEINS USING RNA-PROTEIN
: TITLE OF INVENTION: FUSIONS
: FILE REFERENCE: 00786/350007
: CURRENT APPLICATION NUMBER: US/09/244,796
: CURRENT FILING DATE: 1999-02-05
: EARLIER APPLICATION NUMBER: 60/035,963
: EARLIER FILING DATE: 1997-01-27
: EARLIER APPLICATION NUMBER: 60/064,491
: EARLIER FILING DATE: 1997-11-06
: EARLIER APPLICATION NUMBER: 09/007,005
: EARLIER FILING DATE: 1998-01-14
: NUMBER OF SEQ ID NOS: 33
: SOFTWARE: FastSeq for Windows Version 4.0
: SEQ ID NO 17
: LENGTH: 289
: TYPE: RNA
: ORGANISM: Artificial Sequence
: FEATURE:
: OTHER INFORMATION: Translation template
: NAME/KEY: misc_feature
: LOCATION: (1)...(289)
: OTHER INFORMATION: n = A,T,C or G
US-09-244-796-17

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Query Match          2.4%; Score 38.8; DB 4; Length 289;
Best Local Similarity 5.3%; Pred. No. 0.31;
Matches 10; Conservative 87; Mismatches 93; Indels 0; Gaps 0;

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QY 894 gctctgttgaaagctgtctaacccctccaagaacggatcgcctgctgtcttg 953
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Db 246 GAAAGCTTAAACGCAAGCAAGCAAGCAAGCAAGCAAGCAAGCAAGCAAGCAAG 187
QY 954 acccggtgcaaggcaagcagcaccacatccacacacacacacacacacacac 1013
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 186 NNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNSY 127
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Db 126 NNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNNSNNSY 67
QY 1074 gcatcgagac 1083
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Db 66 NNNNSNNNNY 57

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RESULT 15
US-09-404-650-1
: Sequence 1, Application US/09404650
: Patent No. 6309858
: GENERAL INFORMATION:
: APPLICANT: Dietrich, Paul S.
: APPLICANT: McGovern, Joseph G.
: TITLE OF INVENTION: T-TYPE CALCIUM CHANNEL VARIANTS; COMPOSITIONS THEREOF;

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: TITLE OF INVENTION: AND USES
: FILE REFERENCE: R0043B-REG sequence listing
: CURRENT APPLICATION NUMBER: US/09/404,650
: CURRENT FILING DATE: 1999-09-23
: NUMBER OF SEQ ID NOS: 12
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 1
: LENGTH: 6816
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: CDS
: LOCATION: (192)..(6716)
US-09-404-650-1

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Best Local Similarity 47.5%; Pred. No. 1;
Matches 115; Conservative 0; Mismatches 127; Indels 0; Gaps 0;

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QY 631 catctcgacacccatgctgagatcaagaaggcctcttctgctgtggtggtggtg 690
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Db 4169 caacttcgaaacacttgccagagctctgattgtccctcttgcctggtggtggtggtg 4228
QY 691 gctggcagccctctatggaagcaagaagccttctggtggtggtggtggtggtg 750
      || | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 4229 ttgggtgaaacatcatgtacacatgactggtggtggtggtggtggtggtggtg 4288
QY 751 tgactcatcctggtgctgacatgctacacaggttccctccctggtggtggtggtg 810
      || || || || || || || || || || || || || || || || || || || || ||
Db 4289 caaccacacccctggaatgctgctgtaacttcatctctctcctgctcactgctc 4348
QY 811 tgaacacataaagatgaagaccttgaaggagatctacacagcctggtggtggtggtg 870
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Db 4349 tggctcaacatggttgggtggtggtggtggtggtggtggtggtggtggtggtggtg 4408
QY 871 gg 872
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Db 4409 gg 4410

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Search completed: October 3, 2002, 16:22:08
Job time: 16966 sec

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Fri Oct 4 09:03:06 2002

us-09-826-581-5.rni

Page 10

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 14:48:55 ; Search time 7316.32 Seconds
(without alignments)
288.886 Million cell updates/sec

Title: US-09-826-581-5_COPY_180_280

Perfect score: 101
Sequence: 1 gggccaaagccttgagatgg.....ggccagctgctgagtcacc 101

Scoring table:
IDENTITY_NUC
Gapox 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: gb_ba:*
2: gb_htg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vl:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htgo_inv:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match Length	ID	Description
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1	101	100.0	1647	6	AX281582	AX281582	Sequence
2	101	100.0	2290	9	HS242977	AJ24977	Homo sapi
3	99.4	98.4	2109	6	AX09776	AX09776	Sequence
4	99.4	98.4	2115	6	AX09802	AX09802	Sequence
5	99.4	98.4	2115	6	AF214519	AF214519	Homo sapi
6	72.2	71.5	1867	6	AX09774	AX09774	Sequence
7	72.2	71.5	1873	6	AF214520	AF214520	Sus scrofa
8	72.2	71.5	1873	6	AX09800	AX09800	Sequence
9	72.2	71.5	2022	6	AX09804	AX09804	Sequence
10	71	70.3	989	6	AX281579	AX281579	Sequence
11	71	70.3	152129	2	AC027416	AC027416	Homo sapi
12	71	70.3	206854	9	AC009974	AC009974	Homo sapi
13	69.4	68.7	196554	2	AC073128	AC073128	Homo sapi
14	45.4	45.0	5888	4	AF214521	AF214521	Sus scrofa
15	42.2	41.8	227724	2	AF336381	AF336381	Sus scrofa
16	33	32.7	135012	8	AP003257	AP003257	Mus muscu
17	32	31.7	160133	2	AC024355	AC024355	Oryza sat
18	32	31.7	160878	2	AC016971	AC016971	Homo sapi
19	32	31.7	162840	2	AC046203	AC046203	Homo sapi
20	32	31.7	191330	9	AC025271	AC025271	Homo sapi
21	32	31.7	192175	2	AL591490	AL591490	Mus muscu
22	31.8	31.5	8670	8	CHAK67	CHAK67	Chlamydomon
23	31.6	31.3	124057	9	AP000770	AP000770	Homo sapi
24	31.6	31.3	129750	2	AC103965	AC103965	Homo sapi
25	31.6	31.3	153201	9	AC006454	AC006454	Homo sapi
26	31.6	31.3	165655	2	AC027605	AC027605	Homo sapi
27	31.6	31.3	166941	2	AC073891	AC073891	Homo sapi
28	31.6	31.3	194101	2	AC090910	AC090910	Homo sapi
29	31.6	31.3	210691	9	AP001891	AP001891	Homo sapi
30	31.4	31.1	121851	2	AC091751	AC091751	Gallus ga
31	31.2	30.9	126380	2	AP000714	AP000714	Homo sapi
32	31.2	30.9	139505	9	HSJ591C20	HSJ591C20	Human DNA
33	31.2	30.9	142746	2	AC093564	AC093564	Homo sapi
34	31.2	30.9	157473	2	AL356977	AL356977	Homo sapi
35	31.2	30.9	159840	2	AP000869	AP000869	Homo sapi
36	31.2	30.9	171902	2	AP000846	AP000846	Homo sapi
37	31.2	30.9	198410	2	AP000831	AP000831	Homo sapi
38	31.2	30.9	199321	9	AP000941	AP000941	Homo sapi
39	31	30.7	31549	9	AC096857	AC096857	Homo sapi
40	31	30.7	78505	2	AC006408	AC006408	Homo sapi
41	31	30.7	123551	2	AC005809	AC005809	Homo sapi
42	31	30.7	191832	2	AC091723	AC091723	Sus scrofa
43	30.8	30.5	7459	1	SC8A2	SC8A2	Streptomy
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45	30.8	30.5	177648	3	LMFP265	LMFP265	Leishmani

ALIGNMENTS

RESULT	1	LOCUS	AX281582	Sequence	5 from Patent WO0177305.	DNA	linear	PAT 02-NOV-2001
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DEFINITION	AX281582	VERSION	AX281582.1	GI:16608833
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REFERENCE
AUTHORS
TITLE
JOURNAL
Andersson, L., Luthman, H. and Marklund, S.
Variants of the human amp-activated protein kinase gamma 3 subunit
Patent: WO 0177305-A 5 18-OCT-2001.
Arefis AB (SE)
location/Qualifiers

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/note="unnamed protein product"
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LSPPAPPKLGMDELKPGAQITMRPMOHTCTDMAATSKLVIFDTMLEIKKAFPA
LVANGVAAPLWDSKOSFVGMILITDFILVLRHYRSPVOIYEIENKHETWRELY
LQGCFKPLVSLSPNDLSLEAVYTLIKNRILHRLPYLDVSGNVHLILHKKLKLFLHIF
GSLPRPSFLYRTIODLGICTFERDLAVLETAPILTALDIFVDRVSALPVNVECGOV
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BASE COUNT 346 a 502 c 462 g 337 t

Query Match 100.0%; Score 101; DB 6; Length 1647;
Best Local Similarity 100.0%; Pred. No. 1.7e-16;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gggccaaagccttgatgagacaaagcagaagtcggtgagaaaggagccaccaggtc 60
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DB 180 GGGCCAAAGCCTTGATGATGACAGCAAGCTCGTGAGAGGAGCGACACAGCTC 239
|||||

OY 61 agggggaaggtcccggtccagcgacgtctgtgagtcacc 101
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DB 240 AGGGGGAAGGTCCCGGTCAGGCGACCTGCTGAGTCCACC 280
|||||

RESULT 2
HSA249977 2290 bp mRNA linear PRI 07-APR-2000
LOCUS Homo sapiens mRNA for AMP-activated protein kinase gamma 3 subunit
DEFINITION (Homo sapiens gamma 3 gene).
ACCESSION AJ249977
VERSION AJ249977.1 GI:6688200
KEYWORDS AMP-activated protein kinase; AMPK gamma 3 gene; gamma 3 subunit.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 2290)
AUTHORS Cheung,P.C., Salt,I.P., Davies,S.P., Hardie,D.G. and Carling,D.
TITLE Characterization of AMP-activated protein kinase gamma-subunit
isoforms and their role in AMP binding
JOURNAL Biochem. J. 346 Pt 3, 659-669 (2000)
MEDLINE 20164049
REFERENCE 2 (bases 1 to 2290)
AUTHORS Carling,D.
TITLE Direct Submission
JOURNAL Submitted (12-OCT-1999) Carling D., Cellular Stress Group, MRC
Clinical Sciences Centre, Hammersmith Hospital, DuCane Road,
London, W12 0NN, UNITED KINGDOM
FEATURES
source location/Qualifiers
1. 2290
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22. 1500
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SSERIRGKRRRAKALRWTRBROKSVGEPPGGGEGPRSPRAESGTEATPEKTPPLAQA
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LVANGVAAPLWDSKOSFVGMILITDFILVLRHYRSPVOIYEIENKHETWRELY
LQGCFKPLVSLSPNDLSLEAVYTLIKNRILHRLPYLDVSGNVHLILHKKLKLFLHIF
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BASE COUNT 501 a 674 c 617 g 498 t

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Best Local Similarity 100.0%; Pred. No. 1.6e-16;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 182 GGGCCAAAGCCTTGATGATGACAGCAAGCTCGTGAGAGGAGCGACACAGCTC 241
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OY 61 agggggaaggtcccggtccagcgacgtctgtgagtcacc 101
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DB 242 AGGGGGAAGGTCCCGGTCAGGCGACCTGCTGAGTCCACC 282
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RESULT 3
AX099776 2109 bp DNA linear PAT 02-APR-2001
LOCUS AX099776
DEFINITION Sequence 3 from Patent W00120003.
ACCESSION AX099776
VERSION AX099776.1 GI:13538810
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 2109)
AUTHORS Andersson,L., Looft,C., Kalm,E., Milan,D., Robic,A.,
Rogel-Gallard,C., Iannuccelli,N., Gellin,J., le Roy,P. and
Chardon,P.
TITLE Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
JOURNAL Patent: WO 0120003-A3 22-MAR-2001.
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
FEATURES
source location/Qualifiers
1. 2109
/organism="Homo sapiens"
/db_xref="taxon:9606"
472. 1389
/note="unnamed protein product"
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BASE COUNT 458 a 621 c 560 g 470 t

Query Match 98.4%; Score 99.4; DB 6; Length 2109;
Best Local Similarity 99.0%; Pred. No. 4.1e-16;
Matches 100; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 gggccaaagccttgatgagacaaagcagaagtcggtgagaaaggagccaccaggtc 60
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DB 80 GGGCCAAAGCCTTGATGATGACAGCAAGCTCGTGAGAGGAGCGACACAGCTC 139
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OY 61 agggggaaggtcccggtccagcgacgtctgtgagtcacc 101
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DB 140 AGGGGGAAGGTCCCGGTCAGGCGACCTGCTGAGTCCACC 180
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RESULT 4

AX099802
LOCUS AX099802 2115 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 29 from Patent WO0120003.
ACCESSION AX099802
VERSION AX099802.1 GI:13538836
KEYWORDS human.
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 2115)
AUTHORS Andersson,L., Looft,C., Kalm,E., Milan,D., Robic,A., Rogel-Galliard,C., Iannuccelli,N., Gellin,J., le Roy,P. and Chardon,P.
TITLE Variants of the gamma chain of ampk, dna sequences encoding the same, and uses thereof
JOURNAL Institut NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ; Andersson, Lelf (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE) ; Location/Qualifiers
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BASE COUNT 460 a 622 c 562 g 471 t
ORIGIN

Query Match 98.4% Score 99.4; DB 6; Length 2115;
Best Local Similarity 99.0% Pred. No. 4,1e-16;
Matches 100; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DB 86 GGGCCAAAGCCTTGAGATGACACAGCAGACGCTGCGAGGAGGAGCCACGAGTIC 145
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QY 61 aaggggaaagtcctccggtccaaagcagcgtcgtgagtcacc 101
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DB 146 AGCGGGAAGCTCCCGCTCCAGCCCAACTGCTGAGTCCACC 186
|||||

RESULT 5
AF214519
LOCUS AF214519 2115 bp mRNA linear PRI 03-JUN-2000
DEFINITION Homo sapiens AMP-activated protein kinase gamma subunit (PRKAG3)
ACCESSION AF214519
VERSION AF214519.1 GI:8215681
KEYWORDS human.
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 2115)
AUTHORS Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Thelander,M., Rogel-Galliard,C., Paul,S., Iannuccelli,N., Rask,L., Ronne,H., Lundstrom,K., Reinsch,N., Gellin,J., Kalm,E., Roy,P.L., Chardon,P. and Andersson,L.
TITLE A mutation in PRKAG3 associated with excess glycogen content in pig skeletal muscle

JOURNAL Science 288 (5469), 1248-1251 (2000)
MEDLINE 20280150
PUBMED 10818001
REFERENCE 2 (bases 1 to 2115)
AUTHORS Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Rogel-Galliard,C., Paul,S., Gellin,J., Lundstrom,K., Reinsch,N., Kalm,E., le Roy,P., Chardon,P. and Andersson,L.
TITLE Direct Submission
JOURNAL Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish University of Agricultural Sciences, BMC box 597, Uppsala 751 24, Sweden
FEATURES
source
Location/Qualifiers
1..2115
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="2p"
/map="2p"
/tissue_type="skeletal muscle"
1..2115
/gene="PRKAG3"
1..1395
/gene="PRKAG3"
/note="AMPK3"
/note="AMPK3"
/codon_start=1
/product="AMP-activated protein kinase gamma subunit"
/protein_id="AA73987.1"
/db_xref="GI:8215682"
/translation="MSFLEQENSSSPSPAVTSSSERIRGRRAKALRMWTROKSVEEG
EPFGGEGRSPRTAEVSTGLEATFPPTPLAQADPAGVPTPTGMDCLPSDCTASAG
SSTDVELATEPRATEAMECELEGLERPALCLSPQAFPRKLDLMDLPRGAQIYM
RMOEHCTYDAMATSSKLVIFDTMLEIKKAFALVANGVRAPLMDSKOSFVGLTI
TDFILVHRVYRSPVLOIYEIEOHKLETWREIYLOGCFKPLVSI PNLSFEAVYTLI
KNRIHRLVLPDVPVSGNVLIHTLHKRLKFLHIFCSLPRPSLYRTIODLGIGTRDL
AVVLETPALITLALDIEVDPRVRSALPVNVECGVGLTSRPVTHLAADQTNHLDMSV
GEALRPTLCLEGVISCPHESLSGEVIDRIAREQVHRLVLVDETHLLGVSLSDILQ
ALVSPADIDALGA"
BASE COUNT 460 a 622 c 562 g 471 t
ORIGIN

Query Match 98.4% Score 99.4; DB 9; Length 2115;
Best Local Similarity 99.0% Pred. No. 4,1e-16;
Matches 100; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 gggccaaagccttgatgagacaaagcagaagtcggtgaggaaggagccaccagctc 60
|||||
DB 86 GGGCCAAAGCCTTGAGATGACACAGCAGACGCTGCGAGGAGGAGCCACGAGTIC 145
|||||
QY 61 aaggggaaagtcctccggtccaaagcagcgtcgtgagtcacc 101
|||||
DB 146 AGCGGGAAGCTCCCGCTCCAGCCCAACTGCTGAGTCCACC 186
|||||

RESULT 6
AX099774
LOCUS AX099774 1867 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 1 from Patent WO0120003.
ACCESSION AX099774
VERSION AX099774.1 GI:13538808
KEYWORDS pig.
SOURCE
ORGANISM Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE 1 (bases 1 to 1867)
AUTHORS Andersson,L., Looft,C., Kalm,E., Milan,D., Robic,A., Rogel-Galliard,C., Iannuccelli,N., Gellin,J., le Roy,P. and Chardon,P.
TITLE Variants of the gamma chain of ampk, dna sequences encoding the same, and uses thereof
JOURNAL Institut NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ; Andersson, Lelf (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)


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Query Match      71.5%: Score 72.2: DB 6: Length 1873;
Best Local Similarity 82.2%: Pred. No. 4.6e-09;
Matches 83: Conservative 0: Mismatches 18: Indels 0: Gaps 0:

OY 1 gggccaaagccttgatgagatggaacgaagcagtcgltggaaggaaggaagccaccagtc 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 86 GGAACAGAGCCTCTACATGACACAGCAGCATGTAGACGAAGGGGGCGCTCCGGGCC 145
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

OY 61 agggggaagtcctccggtccagggcagctgctgagtcacc 101
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 146 CGAGGGAAGTCCCGACGTCAGCCACTGCTGAGTCCACC 186
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 9
LOCUS AX099804 2022 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 31 from Patent WO0120003.
ACCESSION AX099804
VERSION AX099804.1 GI:13538838
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
1 (bases 1 to 2022)
Andersson, L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Rogel-Galliard, C., Iannuccelli, N., Gellin, J., Le Roy, P. and
Chardon, P.
TITLE
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 31 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
FEATURES
Location/Qualifiers
1..2022
/organism="Sus scrofa"
/db_xref="taxon:9823"
BASE COUNT 412 a 623 c 593 g 394 t
ORIGIN

Query Match      71.5%: Score 72.2: DB 6: Length 2022;
Best Local Similarity 82.2%: Pred. No. 4.5e-09;
Matches 83: Conservative 0: Mismatches 18: Indels 0: Gaps 0:

OY 1 gggccaaagccttgatgagatggaacgaagcagtcgltggaaggaaggaagccaccagtc 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 236 GGAACAGAGCCTCTACATGACACAGCAGCATGTAGACGAAGGGGGCGCTCCGGGCC 295
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OY 61 agggggaagtcctccggtccagggcagctgctgagtcacc 101
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 296 CGAGGGAAGTCCCGACGTCAGCCACTGCTGAGTCCACC 336
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 10
LOCUS AX281579 989 bp DNA linear PAT 02-NOV-2001
DEFINITION Sequence 2 from Patent WO0177305.
ACCESSION AX281579
VERSION AX281579.1 GI:16608830
KEYWORDS
SOURCE
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
1 (sites)
Andersson, L., Luthman, H. and Marklund, S.
TITLE
Variants of the human amp-activated protein kinase gamma 3 subunit
Patient: WO 0177305-A 2 18-OCT-2001;
JOURNAL
Aresis AB (SE)
FEATURES
Location/Qualifiers
1..989
/organism="Homo sapiens"

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BASE COUNT      229 a 306 c 286 g 168 t
ORIGIN

Query Match      70.3%: Score 71: DB 6: Length 989;
Best Local Similarity 100.0%: Pred. No. 1.1e-08;
Matches 71: Conservative 0: Mismatches 0: Indels 0: Gaps 0:

OY 1 gggccaaagccttgatgagatggaacgaagcagtcgltggaaggaaggaagccaccagtc 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 109 GGGCCAAAGCCTTGACATGACAAAGCAGATCGGTGACGAGGACGACACAGGTC 168
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

OY 61 agggggaagtc 71
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 169 AGGGGGAAGCT 179
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 11
LOCUS AC027416/c 152129 bp DNA linear HTG 07-JUN-2000
DEFINITION Homo sapiens clone RP11-504G11, WORKING DRAFT SEQUENCE. 32
ACCESSION AC027416
VERSION AC027416.2 GI:8317289
KEYWORDS
HTG: HTGS_PHASE1; HTGS_DRAFT.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
1 (bases 1 to 152129)
Birren, B., Linton, L., Nusbaum, C. and Lander, E.
Homo sapiens, clone RP11-504G11
Unpublished
2 (bases 1 to 152129)
Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
Anderson, S., Baldwin, J., Berna, N., Bastien, V., Beda, F.,
Boguslavskiy, L., Boukhgalter, B., Brown, A., Burkett, G.,
Campopiano, A., Castle, A., Choquet, J., Colangelo, M., Collins, S.,
Collins, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S.,
Dodge, S., Domino, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D.,
Galagan, J., Gardyna, S., Ginde, S., Coyette, M., Graham, L.,
Grand-Pierre, N., Grant, G., Hagos, B., Harford, A., Horton, L.,
Howland, J. C., Iliev, I., Johnson, R., Jones, C., Kann, L., Karatas, A.,
Klein, J., Lacombe, K., Lamazares, R., Landers, T., Lehotzky, J.,
Levine, R., Lieu, C., Liu, G., Locke, K., MacDonald, P., Margulis, N.,
McCarthy, M., McEwan, P., McGurk, A., McKernan, K., McPheeters, R.,
Meidirm, J., Menes, L., Mihova, T., Miranda, C., Mlenga, V., Morrow, J.,
Murphy, T., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P.,
O'Neill, D., Oliver, T. M., Oliver, J., Peterson, K., Pierre, N.,
Pisanil, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rotman, D.,
Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B.,
Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,
Tessier, S., Theodore, J., Tirelli, A., Travers, M., Triggillo, J.,
Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J.,
Young, G., Zainoun, J., Zimmer, A. and Zody, M.
Direct Submission
Submitted (30-MAR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 7, 2000 this sequence version replaced gi:7342115.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
-----
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: W1BR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
-----
Project Information
Center project name: L7458
Center clone name: 504_G_11
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Summary Statistics
Sequencing vector: M13; M77815; 100% of reads

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/note="assembly_fragment"

Query Match 70.3%: Score 71: DB 2: Length 152129:

Best Local Similarity 100.0%: Pred. No. 3.8e-09:

Matches 71: Conservative 0: Mismatches 0: Indels 0: Gaps 0:

Oy 1 999ccaaagccttgatgacagaagtcggtgaggaaggagccacagtc 60

Db 2945 GGCCAAAGCCTTGAGATGGACAGCAAGTCGGTGGAGGAAGGAGCCAGCAGTC 2886

Oy 61 aaggaggaaagtc 71

Db 2885 ACGCGCAAGCT 2875

RESULT 12

AC009974/c 206854 bp DNA linear PRI 09-JAN-2002

LOCUS Homo sapiens BAC clone RP11-459119 from 2, complete sequence.

AC009974

AC009974

AC009974.9 GI:16799058

HTG.

KEYWORDS human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

1 (bases 1 to 206854)

Toward a complete human genome sequence

Genome Res. 8 (11), 1097-1108 (1998)

99053792

2 (bases 1 to 206854)

Harris, A. and Colton, M.

The sequence of Homo sapiens BAC clone RP11-459119

Unpublished (2001)

3 (bases 1 to 206854)

Waterston, R.H.

Direct Submission

Submitted (08-SEP-1999) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis,

MO 63108, USA

4 (bases 1 to 206854)

Waterston, R.H.

Direct Submission

Submitted (03-JAN-2002) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis,

MO 63108, USA

5 (bases 1 to 206854)

Waterston, R.H.

Direct Submission

Submitted (09-JAN-2002) Department of Genetics, Washington

University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

On Nov 8, 2001 this sequence version replaced gi:13431203.

----- Genome Center

Center: Washington University Genome Sequencing Center

Center code: WUGSC

Web site: http://genome.wustl.edu/gsc

Contact: saplens@wustl.wustl.edu

----- Summary Statistics

Center project name: H_NH0459119

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RP11-459119 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P. Y., Zhao, B., Frengen, E., Tateo, M., Catanese, J. J., and de Jong, P. J. (1996) An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-1077K22: the clone sequenced to the right is RP11-64705. Actual start of this clone is at base position 1 of RP11-459119; actual end is at base position 206854 of RP11-459119.

Data from AC079810 and AC073128 was used to finish this clone, AC009974. Polymorphisms have been identified between AC073128 and AC009974. A single plasmid region exists between 38812-38903. An unresolved tandem in the HERV SVA exists between 184390-185163. PCR suggests that approximately 1700 bps are missing.

FEATURES

source

1..206854

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/map="2"

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/clone_id="RP11-459119"

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1..37

/note="match to EST BI059713 (NID:g14467240)"

1..37

/note="match to EST BF183086 (NID:g11061273)"

1..37

/note="match to EST AL567345 (NID:g12920610)"

1..37

/note="match to EST AW880850 (NID:g8042860)"

1..37

/note="match to EST BF304755 (NID:g11251653)"

1..37

/note="similar to Homo sapiens EST B1114348 (NID:g14565249)"

1..37

/note="match to EST BG477625 (NID:g13409904)"

1..37

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3..37

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4..37

/note="match to EST A1670836 (NID:g4850567) wa04g10.x1"

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/note="match to EST BG470047 (NID:g13402322)"

281..344

/note="match to EST BE047599 (NID:g8364652) tz39c01.y1"

281..344

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

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misc_feature 281..344 /note="similar to Homo sapiens EST B1114348 (NID:g14565249)"
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294..344 /note="match to EST BG470047 (NID:g13402322)"
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misc_feature /note="match to EST AA043371 (NID:g1521226) zk53e10.r1"
588..929 /note="match to EST T64073 (NID:g667938) yc05d12.r1"
misc_feature 594..763 /note="match to EST BF304755 (NID:g11251653)"
misc_feature 594..763 /note="match to EST BG470047 (NID:g13402322)"
misc_feature 594..763 /note="match to EST AW808050 (NID:g8042860)"
misc_feature 594..731 /note="match to EST BE314060 (NID:g9134719)"
misc_feature 594..764 /note="match to EST B1059713 (NID:g14467240)"
misc_feature 594..763 /note="match to EST BG477625 (NID:g13409904)"
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misc_feature 594..763 /note="match to EST BE908408 (NID:g10402954)"
misc_feature 594..763 /note="match to EST AA043371 (NID:g1521226) zk53e10.r1"
misc_feature 594..763 /note="match to EST A1670836 (NID:g4850567) wa04g10.x1"
misc_feature 594..763 /note="match to EST BF183086 (NID:g11061273)"
misc_feature 594..763 /note="similar to Homo sapiens EST B1114348 (NID:g14565249)"
599..763 /note="match to EST AL567345 (NID:g12920610)"
617..1084 /note="match to EST AA481361 (NID:g2210913) zv44e01.r1"
622..763 /note="match to EST A1860958 (NID:g5514574) w156f05.x1"
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622..763 /note="similar to Mus musculus EST A1196847 (NID:g3749453) u167e01.x1"
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967..1084 /note="match to EST BE908408 (NID:g10402954)"
967..1085 /note="match to EST BF304755 (NID:g11251653)"
967..1091 /note="match to EST AA043371 (NID:g1521226) zk53e10.r1"
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Query Match 70.3% Score 71 DB 9 Length 206854

Best Local Similarity 100.0% Pred. No. 3.6e-09;
Matches 71; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 gggccaaagcccttgagatgacaaagcagaagtgcgttgaggagggggccaccagcgc 60
|||||
Db 168897 GGCCCAAGCCTTGACATGACACAGCAGCAACTCGCTGAGGAAGGAGCCACGAGTC 168838
Qy 61 aggggggaagt 71
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Db 168837 AGGGGAGAGGT 168827
RESULT 13
AC073128 196554 bp DNA linear HTG 21-FEB-2001
AC073128/C Homo sapiens chromosome 2 clone RP11-64705, WORKING DRAFT SEQUENCE.
LOCUS 17 uncloned pieces.
DEFINITION
AC073128.3 GI:13027579
VERSION
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE
ORGANISM
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 196554)
Waterston,R.H.
The sequence of Homo sapiens clone
unpublished
2 (bases 1 to 196554)
Waterston,R.H.
Direct Submission
Submitted (08-JUN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Feb 21, 2001 this sequence version replaced g1:8469048.
COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_NH0647005
Summary Statistics
Sequencing vector: M13; 98%
Sequencing vector: plasmid; 0%
Chemistry: Dye-primer ET; 98% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990119
Consensus quality: 187795 bases at least 040
Consensus quality: 190513 bases at least 030
Insert size: 200000; agarose-fp
Insert size: 194954; sum-of-contigs
Quality coverage: 5.58 in Q20 bases; sum-of-contigs
Quality coverage: 5.67 in Q20 bases; sum-of-contigs
NOTE: This is a 'working draft' sequence. It currently
* consists of 17 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 1157: contig of 1157 bp in length
* 1158 1257: gap of unknown length
* 1258 3600: contig of 2343 bp in length
* 3601 3700: gap of unknown length
* 3701 5103: contig of 1403 bp in length
* 5104 5203: gap of unknown length
* 5204 8524: contig of 3321 bp in length
* 8525 8624: gap of unknown length
* 8625 11856: contig of 3232 bp in length

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* 11957 11956: gap of unknown length
* 11957 15783: contig of 3827 bp in length
* 15784 15883: gap of unknown length
* 15884 21906: contig of 6023 bp in length
* 21907 22006: gap of unknown length
* 22007 28887: contig of 6881 bp in length
* 28888 35255: gap of unknown length
* 35256 35355: gap of unknown length
* 35356 44642: contig of 9287 bp in length
* 44643 44743: gap of unknown length
* 44743 58375: contig of 13533 bp in length
* 58376 58376: gap of unknown length
* 58376 73816: contig of 15441 bp in length
* 73817 73916: gap of unknown length
* 73917 92140: contig of 18224 bp in length
* 92141 92240: gap of unknown length
* 92241 113337: contig of 21097 bp in length
* 113338 113437: gap of unknown length
* 113438 130325: contig of 16888 bp in length
* 130326 130425: gap of unknown length
* 130426 149287: contig of 18862 bp in length
* 149288 149387: gap of unknown length
* 149388 196554: contig of 47167 bp in length.
FEATURES
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Location/Qualifiers
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/db_xref="taxon:9606"
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3701. 5103
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5204. 8524
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8625. 11856
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11957. 15783
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15884. 21906
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22007. 28887
/feature="assembly_name:Contig24"
28988. 35255
/feature="assembly_name:Contig25"
35356. 44642
/feature="assembly_name:Contig26"
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/feature="assembly_name:Contig27"
58376. 73816
/feature="assembly_name:Contig28"
73917. 92140
/feature="assembly_name:Contig29"
92241. 113337
/feature="assembly_name:Contig30"
113438. 130325
/feature="assembly_name:Contig31"
130426. 149287
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149388. 196554
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ORIGIN
Query Match 68.7%: Score 69.4: DB 2: length 196554:
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Best Local Similarity 98.6%: Pred. No. 9.4e-09:
Matches 70: Conservative 0: Mismatches 1: Indels 0: Gaps 0:
Oy 1 gggccaaagccttgatgacagaagtcggtggaaggaagggagaccacagtc 60
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Db 63318 GGCCAAAGCCTTGAGATGACAAAGCAGAAAGTCGCGAGCAAGGAGGACACAGATC 63259
|||||
Oy 61 agggggaaggt 71
|||||
Db 63258 AGGGGGAAGGT 63248

RESULT 14
AF214521 5888 bp DNA linear MAM 03-JUN-2000
LOCUS
DEFINITION
ACCESSION
AF214521
VERSION
AF214521.1 GI:8215685
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Euteria; Cetartiodactyla; Suina; Suidae; Sus.
1 (bases 1 to 5888)
Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Thelander,M.,
Rogel-Gallard,C., Paul,S., Iannuccelli,N., Raak,L., Ronne,H.,
Lundstrom,K., Reinsch,N., Gellin,J., Kalm,E., Roy,P.L., Chardon,P.
and Anderson,L.
A mutation in PRAG3 associated with excess glycogen content in pig
skeletal muscle
Science 288 (5469), 1248-1251 (2000)
20280150
PUBMED
10818001
2 (bases 1 to 5888)
Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A.,
Rogel-Gallard,C., Paul,S., Gellin,J., Lundstrom,K.,
Kalm,E., Le Roy,P., Chardon,P. and Andersson,L.
Direct Submission
Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish
University of Agricultural Sciences, BMC Box 597, Uppsala 751 24,
Sweden
FEATURES
source
Location/Qualifiers
1. 5888
/organism="Sus scrofa"
/db_xref="taxon:9823"
/chromosome="15"
/map="15q"
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2771.2825,3027.3153,3286.3451,4578.4615,4791.4937,
5294.5410)
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/product="AMPK gamma subunit"
<1.5410
/feature="PRAG3"
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5294.5410)
/feature="PRAG3"
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/product="AMPK gamma subunit"
/protein_id="AAF33989.1"
/db_xref="GI:8215686"
/translation="MSFLDEGESRSPRAVTTSSERSHQDNKASRWTRQEDVEG
GPGPREGQSRPVAESTGQDAFPKATPLAQAPLAEDVNDPTERRDILPDCASAS
DSNTDHLDLGIIEFSASASGDELGLVEERKAPCPSPPEVLPRGMDDELPGCAQYV
HFMQERTCYDAMATSSKLYIEDTMLIEKKAFVLVANGVRAAPLMDSKOSPGMGLTI
TDFILVLRHYRSPILVOIYEIEBEKLEETMEETIYLGCCFPRAVVISPNDSLFPAVYALI
KNRIHLRPVLDPVSGAVILHTPKRLKEHLHITGTLIPRPSFLIRTTDLGIGTGRDL
AVYLERAPLITLADITFVDRVSAALPVYNETGOVGLYSRFDVTHLAAGQTVHRLDMNV
GEALRORTLCLEGVLSQPHETLTGEVIDRIVROVRLVDETQHLGLCVSLSDTLQ
ALVLSAPGIDALCA"
BASE COUNT 1274 a 1634 c 1638 g 1339 t 3 others
ORIGIN
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Fri Oct 4 09:03:08 2002

us-09-826-581-5_copy_180_280.rge

Page 11

PN W0200177305-A2.
XX 18-OCT-2001.
XX 06-APR-2001; 2001WO-SE00765.
XX 07-APR-2000; 2000US-195665P.
XX (AREX-) AREXIS AB.
XX Andersson L, Luthman H, Marklund S;
PI WPI: 2001-657170/75.
DR P-PSDB: Q0B47679.
XX
XX New variants of human AMP-activated protein kinase gamma3 subunit
PT associated with a metabolic disease e.g. diabetes or obesity and method
PT for determining a risk estimate of diseases in subject by detecting the
PT variant.
XX
XX Disclosure: Fig 5; 25pp; English.
XX
XX This sequence represents the full length cDNA encoding the human
CC AMP-activated protein kinase gamma 3 subunit (PRKAG3). Detecting
CC the presence of the PRKAG3 DNA, or a variant, is useful in determining
CC a risk estimate of a metabolic disease, such as diabetes or obesity,
CC in a subject. The variation may occur in exons 3, 4 or 10. In exon
CC 3 a variation may be a substitution of a G for a C at nucleotide 320,
CC resulting in the amino acid substitution P71A; in exon 4 variation may
CC be a substitution of a T for a C at nucleotide 550; and in exon 10
CC variation may be a substitution of a T for a C at nucleotide 1037,
CC resulting in the amino acid substitution R340W. There may also be
CC nucleotide variation in intron 6. The numbering of these
CC variations is based on the full length cDNA as given, rather than on
CC position 1 of the open reading frame.
XX
XX Sequence 1647 BP; 346 A; 502 C; 462 G; 337 T; 0 other;
SQ

Query Match 100.0%; Score 101; DB 22; Length 1647;
Best Local Similarity 100.0%; Pred. No. 1.3e-20;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gggccaagccttgagatgagcaagaagtcggtgaggaaggagccaccaggtc 60
DB 180 gggccaagccttgagatgagcaagaagtcggtgaggaaggagccaccaggtc 239
OY 61 agggggaaggtcccggtccagcgagcagctgctgagtcacc 101
DB 240 agggggaaggtcccggtccagcgagcagctgctgagtcacc 280

RESULT 2
AAD03296
ID AAD03296 standard; DNA; 2109 BP.
XX
XX AAD03296:
XX 13-JUN-2001 (first entry)
XX
XX Human AMK gamma subunit muscle-specific isoform, PRKAG3 cDNA.
DE
XX Human; gamma subunit; adenosine monophosphate-activated kinase; AMPK;
KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
KW cystathione beta synthase; CBS; cardiac; gene therapy; ss.
XX
XX Homo sapiens.
OS
XX
XX Key Location/Qualifiers
FH 1..471
FT /*tag- a
FT 472..1389
FT CDS

FT /*tag- b
FT /product- "Human Prkag3 protein"
FT 3'UTR 1390..2109
FT /*tag- c
XX
XX W0200120003-A2.
XX 22-MAR-2001.
XX 11-SEP-2000; 2000WO-EP09896.
XX 10-SEP-1999; 99EP-0402236.
XX 18-MAY-2000; 2000EP-0401388.
XX
XX (INRG) INRA INST NAT RECH AGRONOMIQUE.
PA (ANDE/) ANDERSSON L.
PA (LOOF/) LOOFF C.
PA (KALM/) KALM E.
XX
XX Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
PI Iannuccielli N, Gellin J, Le Roy P, Chardon P;
DR WPI: 2001-244810/25.
DR P-PSDB: AAE00221.
XX
XX New variants of the gamma subunit of vertebrate adenosine
PT monophosphate-activated kinase for diagnosis or treatment of disorders
PT associated with energy metabolism such as diabetes, obesity, and
PT myopathy.
XX
XX Claim 12: Fig 2; 71pp; English.
XX
XX The present sequence is a cDNA encoding human adenosine monophosphate
CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
CC PRKAG3. Mutation in Prkag3 results in an altered regulation of
CC carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
CC useful as therapeutic for treating carbohydrate metabolism disorders such
CC as diabetes, obesity, and disorders associated with muscle metabolism
CC such as myopathy and cardiovascular diseases, to modulate AMPK
CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
CC and its functionally altered mutants are useful for the diagnostic
CC evaluation, genetic testing and prognosis of a metabolic disorder.
CC preferably a carbohydrate metabolism disorder. Primers that can detect
CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
CC useful for detecting a dysfunction of carbohydrate metabolism resulting
CC from the expression of a functionally altered allele of PRKAG3.
CC Transgenic animal and host cell transformed with PRKAG3 or a
CC heterotimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC screening compounds able to modulate AMPK activity. Nucleic acid
CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
CC in a sequence encoding the first cystathione beta synthase (CBS) domain
CC of PRKAG3 and is useful in gene therapy.
XX
XX Sequence 2109 BP; 458 A; 621 C; 560 G; 470 T; 0 other;
SQ

Query Match 98.4%; Score 99.4; DB 22; Length 2109;
Best Local Similarity 99.0%; Pred. No. 4.1e-20;
Matches 100; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 gggccaagccttgagatgagcaagaagtcggtgaggaaggagccaccaggtc 60
DB 80 gggccaagccttgagatgagcaagaagtcggtgaggaaggagccaccaggtc 139
OY 61 agggggaaggtcccggtccagcgagcagctgctgagtcacc 101
DB 140 agggggaaggtcccggtccagcgagcagctgctgagtcacc 180

RESULT 3
AAD03320
ID AAD03320 standard; cDNA; 2115 BP.
XX

AC AAD0320;
 XX
 DT 13-JUN-2001 (first entry)
 XX
 DE Human AMPK gamma subunit muscle-specific isoform, complete PRKAG3 cDNA.
 XX
 KW Human: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
 KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
 KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
 KW cystathione beta synthase; CBS; cardiant; gene therapy; ss.
 XX
 OS Homo sapiens.
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 FH Key Location/Qualifiers
 FT 1..1395
 FT CDS /*tag= a
 FT /product= "Human complete Prkag3 protein"
 XX
 PN W0200120003-A2.
 XX
 PD 22-MAR-2001.
 XX
 PP 11-SEP-2000; 2000WO-EP09896.
 XX
 PR 10-SEP-1999; 99EP-0402236.
 PR 18-MAY-2000; 2000EP-0401388.
 XX
 PA (INRG) INRA INST NAT RECH AGRONOMIQUE.
 PA (ANDE/) ANDERSSON L.
 PA (LOOFT/) LOOFT C.
 PA (KALM/) KALM E.
 XX
 PI Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
 PI Iannuccielli N, Gellin J, Le Roy P, Chardon P;
 XX
 DR WPI: 2001-244810/25.
 DR P-PSDB: AAE00223.
 XX
 DT New variants of the gamma subunit of vertebrate adenosine
 DT monophosphate-activated kinase for diagnosis or treatment of disorders
 DT associated with energy metabolism such as diabetes, obesity, and
 DT myopathy -
 XX
 PS Claim 12: Page 65-68; 71pp: English.
 PS
 XX
 CC The present sequence is a cDNA encoding human adenosine monophosphate
 CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
 CC complete PRKAG3. Mutation in Prkag3 results in an altered regulation of
 CC carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
 CC useful as therapeutic for treating carbohydrate metabolism disorders such
 CC as diabetes, obesity, and disorders associated with muscle metabolism
 CC such as myopathy and cardiovascular diseases, to modulate AMPK
 CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
 CC and its functionally altered mutants are useful for the diagnostic
 CC evaluation, genetic testing and prognosis of a metabolic disorder,
 CC preferably a carbohydrate metabolism disorder. Primers that can detect
 CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
 CC useful for detecting a dysfunction of carbohydrate metabolism resulting
 CC from the expression of a functionally altered allele of PRKAG3.
 CC Transgenic animal and host cell transformed with PRKAG3 or a
 CC heterotimeric AMPK consisting of PRKAG3 or its mutant, are useful for
 CC screening compounds able to modulate AMPK activity. Nucleic acid
 CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
 CC in a sequence encoding the first cystathione beta synthase (CBS) domain
 CC of PRKAG3 and is useful in gene therapy.
 XX
 SO Sequence 2115 BP; 460 A; 622 C; 562 G; 471 T; 0 other;

Query Match 98.4%: Score 99.4; DB 22; Length 2115;
 Best Local Similarity 99.0%: Pred. No. 4; le-20;
 Matches 100; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 gggccaaagccttgatgacgaagcagaagtcgtgaggaagggaagccaccagtc 60
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 86 gggccaaagccttgatgacgaagcagaagtcgtgaggaagggaagccaccagtc 145
 OY 61 aggggggaaggtcccggtccaggcagctgctgctcacc 101
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 146 aggggggaaggtcccggtccaggcagctgctgctcacc 186
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 AAD03295
 ID AAD03295 standard; CDNA: 1867 BP.
 XX
 AC AAD03295;
 XX
 DT 13-JUN-2001 (first entry)
 DT
 XX
 DE Pig AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.
 XX
 KW Pig: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
 KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
 KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
 KW cystathione beta synthase; CBS; cardiant; gene therapy; RN locus;
 KW chromosome 15; ss.
 XX
 OS Sus scrofa.
 XX
 FH Key Location/Qualifiers
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 FT 5'UTR
 FT CDS /*tag= a
 FT 472..1389
 FT /*tag= b
 FT /product= "Sus scrofa PRKAG3 protein"
 FT 1390..1867
 FT /*tag= c
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 PN W0200120003-A2.
 XX
 PD 22-MAR-2001.
 XX
 PP 11-SEP-2000; 2000WO-EP09896.
 XX
 PR 10-SEP-1999; 99EP-0402236.
 PR 18-MAY-2000; 2000EP-0401388.
 XX
 PA (INRG) INRA INST NAT RECH AGRONOMIQUE.
 PA (ANDE/) ANDERSSON L.
 PA (LOOFT/) LOOFT C.
 PA (KALM/) KALM E.
 XX
 PI Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
 PI Iannuccielli N, Gellin J, Le Roy P, Chardon P;
 XX
 DR WPI: 2001-244810/25.
 DR P-PSDB: AAE00220.
 XX
 DT New variants of the gamma subunit of vertebrate adenosine
 DT monophosphate-activated kinase for diagnosis or treatment of disorders
 DT associated with energy metabolism such as diabetes, obesity, and
 DT myopathy -
 XX
 PS Claim 12: Fig 2: 71pp: English.
 PS
 XX
 CC The present sequence is a cDNA encoding pig adenosine monophosphate
 CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
 CC complete PRKAG3. Prkag3 gene is located in the RN locus of chromosome 15.
 CC Mutation in Prkag3 results in an altered regulation of carbohydrate
 CC metabolism, particularly in skeletal muscle. PRKAG3 is useful as
 CC therapeutic for treating carbohydrate metabolism disorders such as
 CC diabetes, obesity, and disorders associated with muscle metabolism
 CC such as myopathy and cardiovascular diseases, to modulate AMPK
 CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
 CC and its functionally altered mutants are useful for the diagnostic


```

XX (INRG ) INRA INST NAT RECH AGRONOMIQUE.
PA (ANDE/) ANDERSSON L.
PA (LOOF/) LOOFT C.
PA (KALM/) KALM E.
XX
PI Andersson L, Loof C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
PI Iannucciell N, Geilin J, Le Roy P, Chardon P;
XX
DR WPI: 2001-244810/25.
DR P-PSDB: AAE00224.
XX
PT New variants of the gamma subunit of vertebrate adenosine
PT monophosphate-activated kinase for diagnosis or treatment of disorders
PT associated with energy metabolism such as diabetes, obesity, and
PT myopathy -
XX
PS Claim 12: Page 69: 71pp: English.
XX
CC The present sequence is pig adenosine monophosphate (AMP)-activated
CC kinase (AMPK) gamma subunit muscle-specific isoform, PRKAG3 splice
CC variant DNA. prkag3 gene is located in the RN locus of chromosome 15.
CC Mutation in Prkag3 results in an altered regulation of carbohydrate
CC metabolism, particularly in skeletal muscle. PRKAG3 is useful as
CC therapeutic for treating carbohydrate metabolism disorders such as
CC diabetes, obesity, and disorders associated with muscle metabolism
CC such as myopathy and cardiovascular diseases, to modulate AMPK
CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
CC and its functionally altered mutants are useful for the diagnostic
CC evaluation, genetic testing and prognosis of a metabolic disorder,
CC preferably a carbohydrate metabolism disorder. Primers that can detect
CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
CC useful for detecting a dysfunction of carbohydrate metabolism resulting
CC from the expression of a functionally altered allele of PRKAG3.
CC Transgenic animal and host cell transformed with PRKAG3 or a
CC heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC screening compounds able to modulate AMPK activity. Nucleic acid
CC encoding PRKAG3 is useful for detecting mutations in a prkag3 gene, or
CC in a sequence encoding the first cystathione beta synthase (CBS) domain
CC of PRKAG3 and is useful in gene therapy.
XX
SQ Sequence 2022 BP: 412 A: 623 C: 593 G: 394 T: 0 other:

```

```

Query Match      71.5%: Score 72.2: DB 22: Length 2022:
Best Local Similarity 82.2%: Pred. No. 4.1e-12:
Matches 83: Conservative 0: Mismatches 18: Indels 0: Gaps 0:
OY 1 gggccaaagccttgagatgacaaagcagaagtcgltgagaaaggagccaccaggtc 60
   || || || || || || || || || || || || || || || || || || || ||
DB 236 ggaacaagcctctgagatgacaaagcagaagtcgltgagaaaggagccaccaggtc 295
OY 61 agggggaaggtcccggtccaggtcgtgagtcacc 101
   | || || || || || || || || || || || || || || || || || || ||
DB 296 cgaagggaaggtcccggtccaggtcgtgagtcacc 336

```

```

RESULT 7
AAH43682 standard: DNA: 989 BP.
ID AAH43682
AC AAH43682:
XX
XX 21-JAN-2002 (first entry)
XX
XX PRKAG3 Intron 2 - Intron 4.
XX
XX Human: AMP-activated protein kinase gamma 3 subunit; PRKAG3; variant;
XX metabolic disease; diabetes; obesity; substitution; ds.
OS Homo sapiens.
XX
XX Key Location/Qualifiers
FH

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FT /number= "Intron 2"
FT /note= "3' portion of Intron 2"
FT 22..177
FT /*tag= b
FT /number= "Exon 3"
FT 178..541
FT /*tag= c
FT /number= "Intron 3"
FT 542..945
FT /*tag= d
FT /number= "Exon 4"
FT 946..989
FT /*tag= e
FT /number= "Intron 4"
FT /note= "5' portion of Intron 4"
XX
XX W0200177305-A2.
XX
XX 18-OCT-2001.
XX
XX 06-APR-2001: 2001WO-SE00765.
XX
XX 07-APR-2000: 2000US-195665P.
XX
XX (AREX-) AREXIS AB.
XX
XX Andersson L, Luthman H, Marklund S;
XX
XX WPI: 2001-657170/75.
XX

```

New variants of human AMP-activated protein kinase gamma3 subunit associated with a metabolic disease e.g. diabetes or obesity and method for determining a risk estimate of diseases in subject by detecting the variant -

Example 1; Fig 2: 25pp: English.

The sequences given in AAH43681-84 represents genomic fragments encoding the human AMP-activated protein kinase gamma 3 subunit (PRKAG3). Detecting the presence of the PRKAG3 DNA, or a variant, is useful in determining a risk estimate of a metabolic disease, such as diabetes or obesity, in a subject. The variation may occur in exons 3, 4 or 10. In exon 3 variation may be a substitution of a G for a C at nucleotide 320, resulting in the amino acid substitution P71A; in exon 4 variation may be a substitution of a T for a C at nucleotide 550; and in exon 10 variation may be a substitution of a T for a C at nucleotide 1037, resulting in the amino acid substitution R340W. There may also be nucleotide variation in intron 6.

Sequence 989 BP: 229 A: 306 C: 286 G: 168 T: 0 other:

```

Query Match      70.3%: Score 71: DB 22: Length 989:
Best Local Similarity 100.0%: Pred. No. 8.4e-12:
Matches 71: Conservative 0: Mismatches 0: Indels 0: Gaps 0:
OY 1 gggccaaagccttgagatgacaaagcagaagtcgltgagaaaggagccaccaggtc 60
   || || || || || || || || || || || || || || || || || || || ||
DB 109 gggccaaagccttgagatgacaaagcagaagtcgltgagaaaggagccaccaggtc 168
OY 61 agggggaaggt 71
   | || || || || || || || || || || || || || || || || || || ||
DB 169 agggggaaggt 179

```

```

RESULT 8
AAZ32059/c
ID AAZ32059 standard: DNA: 8670 BP.
XX
XX AAZ32059:
AC

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Query Match 31.5%; Score 31.8; DB 22; Length 8670;
Best Local Similarity 64.0%; Pred. No. 3.8;
Matches 48; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

QY 14 gaaatggaacaaagcagagatcgtaggaagaaggaccacagtcaggagggaagc 73
DB 6582 GAGCTGGACGACTGGCAGACCCGCTGACGAGCTGGAGGCGGATCATCAACAGCCG 6523
OY 74 ccggtccaggccagc 88
DB 6522 CCGGCCCAACCCCGC 6508

RESULT 10
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ID AAS84420 standard; cDNA: 711 BP.
XX
AC AAS84420;
XX
UT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #20224.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
HU 11-OCT-2001.
XX
PP 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
DR P-PSDB: ABG20233.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity.
XX
PS Claim 1: SEQ ID No 20224; 103pp: English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. AAS64197-AAS94564 represent novel human
XX diagnostic coding sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO

CC at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX Sequence 711 BP; 139 A; 227 C; 249 G; 96 T; 0 other;
S0

Query Match 29.7%; Score 30; DB 23; Length 711;
Best Local Similarity 67.7%; Pred. No. 9.2;
Matches 42; Conservative 0; Mismatches 20; Indels 0; Gaps 0;

QY 30 aatcggtggaagaaggagccacacagtcaggagggaagtcctccggtccagccagct 89
DB 431 aactcgctcggaagaagagacaccccggggcaagaagaaacctccagcgaagaagactgct 490
OY 90 gc 91
DB 491 gc 492

RESULT 11
AAS84523
ID AAS84523 standard; cDNA: 2541 BP.
XX
AC AAS84523;
XX
UT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #20327.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
HU 11-OCT-2001.
XX
PP 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
DR P-PSDB: ABG20336.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity.
XX
PS Claim 1: SEQ ID No 20327; 103pp: English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. AAS64197-AAS94564 represent novel human

CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_ptc_sequences.
XX
SQ Sequence 2541 BP; 476 A; 780 C; 787 G; 498 T; 0 other;

Query Match 29.5%; Score 29.8; DB 23; Length 2541;
Best Local Similarity 63.0%; Pred. No. 13;
Matches 46; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

Qy 15 agatggacaagcgaagtcggtgaggaagcaccaggtcagggaagctcc 74
Db 1104 agagccggtgcaaggcggcggtggaacttgcggccgaggtcagggtctc 1163
Oy 75 cgtccagccag 87
Db 1164 cagtcaccgctt 1176

RESULT 12

ID AAD22004 standard; CDNA: 4125 BP.

AC AAD22004;

DT 12-FEB-2002 (first entry)

DE Human transporters and ion channels (TRICH)-12 CDNA.

XX Human; transporter and ion channel; TRICH: akinesia; cystic fibrosis;
KM diabetes mellitus; Parkinson's disease; myasthenia gravis; dementia;
KM cardiac disorder; angina; hypertension; myocarditis; hyperglycemia;
KM neurological disorder; Alzheimer's disease; cataract; infertility;
KM Wilson's disease; schizophrenia; Grave's disease; Addison's disease;
KM Huntington's disease; multiple sclerosis; meningitis; hypotensive;
KM cardiac; nocturnal; neuroprotective; neuroleptic; ophthalmological;
KM antihypertensive; anticonvulsant; goitre; antiinflammatory; ss.

XX Homo sapiens.

XX Key Location/Qualifiers
FT CDS 1..3933

FT /*tag= a
FT /product= "Human transporters and ion channels
FT (TRICH)-12"

PN MO200177174-A2.

XX 18-OCT-2001.

PD 06-APR-2001; 2001MO-US11206.

XX 06-APR-2001; 2001MO-US11206.

XX 06-APR-2001; 2000US-195595P.

PR 12-APR-2000; 2000US-196872P.

PK 20-APR-2000; 2000US-199020P.

PR 28-APR-2000; 2000US-200552P.

PR 05-MAY-2000; 2000US-202348P.

PR 11-MAY-2000; 2000US-203495P.

XX (INCY-) INCYTE GENOMICS INC.

PA (INCY-) INCYTE GENOMICS INC.

XX Reddy R, Thornton M, Borowsky ML, Tang YT, Khan FA, Tribouley CM;

PI Gandhi AR, Yao MG, Sanjwal MS, Baughn MR, Nguyen DB;

PI Policky JL, Yue H, Selthamer JJ, Walla NK, Lal P, Kearney L;

PI Walsh RT, Lu DAM, Lu Y, Greene BD, Raumann BE, Patterson C;

XX WPI: 2002-017448/02.

DR P-PSDB: AAE13285.

XX Polypeptides of human transporters and ion channels, useful for

PT diagnosing, treating or preventing disorders of transport.

PT neurological, muscle, immunological and cell proliferative disorders -
XX
PS Claim 5; Page 147-148; 150pp; English.

CC The invention relates to human transporters and ion channels (TRICH)
CC and the polynucleotides encoding them. The composition comprising TRICH
CC or agonist of TRICH is useful for treating a disease or condition
CC associated with decreased expression of functional TRICH or condition
CC associated with overexpression of TRICH respectively. The composition
CC comprising Ab is useful for diagnosing a condition of disease associated
CC with expression of TRICH in a subject, where the disorders include a
CC transport disorder such as akinesia, cystic fibrosis, diabetes mellitus,
CC Parkinson's disease, myasthenia gravis, cardiac disorders associated
CC with transport e.g. angina, hypertension, myocarditis, neurological
CC disorders associated with transport e.g. Alzheimer's disease, Wilson's
CC disease, schizophrenia, cataracts, infertility, hyperglycemia, Grave's
CC disease, goitre, Addison's disease, Huntington's disease, dementia,
CC multiple sclerosis, bacterial and viral meningitis. TRICH DNA is useful
CC for generating a transcript image of a tissue or cell type, which
CC represents the global pattern of gene expression by a particular tissue
CC or cell type and for analysing the proteome of a tissue or cell type.
CC TRICH DNA is used in gene therapy. The present sequence is human
CC TRICH12 CDNA.

SQ Sequence 4125 BP; 989 A; 1109 C; 1188 G; 839 T; 0 other;

Query Match 29.5%; Score 29.8; DB 24; Length 4125;
Best Local Similarity 60.5%; Pred. No. 13;

Matches 49; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

Oy 7 aagccttgatgacagcagcagcagcagcagcagcagcagcagcagcagc 66
Db 3901 aagccttgatgacagcagcagcagcagcagcagcagcagcagcagc 3960
Oy 67 aaggtcccggtcagcagcagcagcagcagcagcagcagcagcagcagc 87
Db 3961 atcctctgagcagcagcagcagcagcagcagcagcagcagcagc 3981

RESULT 13

AAH06841/C
ID AAH06841 standard; CDNA: 886 BP.

AC AAH06841;

DT 26-JUN-2001 (first entry)

DE Human CDNA clone (5'-primer) SEQ ID NO:3676.

XX Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

XX Homo sapiens.

PN EP1074617-A2.

PD 07-FEB-2001.

PF 28-JUL-2000; 2000EP-0116126.

PR 29-JUL-1999; 99JP-0248036.

PR 27-AUG-1999; 99JP-0300253.

PR 11-JAN-2000; 2000JP-0118776.

PR 02-MAY-2000; 2000JP-0183767.

PR 09-JUN-2000; 2000JP-0241899.

XX (HELI-) HELIX RES INST.

PA (HELI-) HELIX RES INST.

XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX WPI: 2001-318749/34.

PT primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs -

XX
XX
PS Claim 1: SEQ ID 3676; 2537pp + CD ROM; English.

CC The present invention describes primer sets for synthesizing 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesizing polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.

XX
SQ Sequence 886 BP; 169 A; 259 C; 213 G; 241 T; 4 other;

Query Match 29.1%; Score 29.4; DB 22; Length 886;
Best Local Similarity 58.6%; Pred. No. 14;
Matches 51: Conservative 0; Mismatches 36; Indels 0; Gaps 0;

OY 2 ggcacaaagccttgatgacaaagcagaagtcgltgaggaaggagccaccagatca 61
DB 100 GCCTAAGGACTCCGTAGGCGCCAGCGCGAGCGCTCGAGAGCAGCAGCGCC 41
OY 62 gggggaaggtcccggtccagcagc 88
DB 40 AGAGAGAACTCCCGGGCCAGCCTGC 14

RESULT 14
AAH14551/C
ID AAH14551 standard; cDNA; 1457 BP.

XX
AC AAH14551;

XX
DT 26-JUN-2001 (first entry)

XX
DE Human cDNA sequence SEQ ID NO:12117.

XX
KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

XX
OS Homo sapiens.

XX
PN EP1074617-A2.

XX
PD 07-FEB-2001.

XX
PE 28-JUL-2000; 2000EP-0116126.

XX
PR 29-JUL-1999; 99JP-0248036.

XX
PR 27-AUG-1999; 99JP-0300253.

XX
PR 11-JAN-2000; 2000JP-0118776.

XX
PR 02-MAY-2000; 2000JP-0183767.

XX
PR 09-JUN-2000; 2000JP-0241899.

XX
PA (HELI-) HELIX RES INST.

XX
XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
XX WPI: 2001-318749/34.

PT primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs -

PS Claim 8: SEQ ID 12117; 2537pp + CD ROM; English.

CC The present invention describes primer sets for synthesizing 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesizing polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.

XX
SQ Sequence 1457 BP; 291 A; 442 C; 365 G; 359 T; 0 other;

Query Match 29.1%; Score 29.4; DB 22; Length 1457;
Best Local Similarity 58.6%; Pred. No. 15;
Matches 51: Conservative 0; Mismatches 36; Indels 0; Gaps 0;

OY 2 ggcacaaagccttgatgacaaagcagaagtcgltgaggaaggagccaccagatca 61
DB 100 GCCTAAGGACTCCGTAGGCGCCAGCGCGAGCGCTCGAGAGCAGCAGCGCC 41
OY 62 gggggaaggtcccggtccagcagc 88
DB 40 AGAGAGAACTCCCGGGCCAGCCTGC 14

RESULT 15
AAV59752/C
ID AAV59752 standard; DNA; 1482 BP.

XX
AC AAV59752;

XX
DT 19-JAN-1999 (first entry)

XX
DE Human secreted protein gene 96 clone HAQBK61.

XX
KW Human; secreted protein; fusion protein; gene therapy; protein therapy;

XX
KW diagnosis; tissue; cancer; tumour; neurodegenerative disorder; leukaemia;

XX
KW developmental abnormality; foetal deficiency; blood; allergy; renal; ds;

XX
KW immune system; asthma; lymphocytic disease; brain; hepatic; lymphoma;

XX
KW inflammation; ischaemic shock; Alzheimer's disease; restenosis; AIDS;

XX
KW cognitive disorder; schizophrenia; prostate; obesity; osteoclast; thymus;

XX
KW osteoporosis; arthritis; testis; lung; thyroiditis; thyroid; digestion;

XX
KW endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.

XX
OS Homo sapiens.

Fri Oct 4 09:03:08 2002

us-09-826-581-5_copy_180_280.rng

Page 11

pb 90 GCGCTAAAGCACTCCGTTAGGGGGCCAGCGCGCAACGGCGGAGCGCTCGGAGGGAGCACAGGGCC 31

OY 62 qgqgaaagyltccccag lcccaagcagc 88

Db 30 ACAGAGAGAGTCCCGCGGGCGCACCTTGC 4

Search completed: October 3, 2002, 16:30:43
Job time: 14313 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:15:01 ; Search time 5701.1 Seconds
(without alignments)
239.110 Million cell updates/sec

Title: US-09-826-581-5_COPY_180_280

Perfect score: 101

Sequence: 1 gggccaaagccttgatgagtg9.....ggccagctgctgagtcacac 101

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 13736207 segs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estlu:*
4: em_estmu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_htc:*
9: gb_estcl:*
10: gb_estcl2:*
11: gb_htc:*
12: gb_gss:*
13: em_gss_hum:*
14: em_gss_luv:*
15: em_gss_pln:*
16: em_gss_vrc:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	67.4	66.7	399	9	AM356079
2	67.4	66.7	422	9	AM427435
3	67.4	66.7	621	9	BB628877
4	67.4	66.7	653	9	BB629521
5	65.8	65.1	685	9	BB630381
6	54.6	54.1	548	10	BI775360
7	53.8	53.3	516	9	AI664508
8	52.6	52.1	444	10	BF890374
9	35.6	35.2	922	10	BI754053
10	32.6	32.3	406	9	BB759831
11	32.6	32.3	457	9	BB780864
12	32.6	32.3	736	10	BI765906
13	32.4	32.1	151	10	BG244994
14	32	31.7	454	9	BB860266
15	32	31.7	479	10	BF549656
16	31.2	30.9	896	10	BF526844
17	31	30.7	327	9	BB753946

18	31	30.7	1074	10	BG966577
19	30.8	30.5	365	9	AA787203
20	30.8	30.5	491	10	BF651383
21	30.8	30.5	722	12	A2188883
22	30.6	30.3	512	9	AI939950
23	30.6	30.3	1195	10	BE299786
24	30.4	30.1	370	10	BG086356
25	30.4	30.1	443	9	BB845787
26	30.4	30.1	514	10	BI181375
27	30.2	29.9	167	10	BF393187
28	30.2	29.9	211	9	BB438733
29	30.2	29.9	428	10	BF394635
30	30.2	29.9	545	9	AM413109
31	30.2	29.9	825	10	BI910794
32	30.2	29.9	1079	12	CNS05J0V
33	30	29.7	250	10	BE426030
34	30	29.7	409	10	BE426831
35	30	29.7	429	10	BE489634
36	30	29.7	1253	11	AK015101
37	29.8	29.5	335	10	225209
38	29.8	29.5	385	12	A0852870
39	29.8	29.5	581	12	BH414079
40	29.8	29.5	662	12	CNS048SD
41	29.8	29.5	740	10	BF135550
42	29.8	29.5	1101	12	CNS00004
43	29.6	29.3	285	9	BB039663
44	29.6	29.3	375	9	AM298840
45	29.6	29.3	431	9	BB822055

ALIGNMENTS

RESULT 1
LOCUS 38073 MARC 2BOV Bos taurus CDNA 5', mRNA sequence.
DEFINITION AM356079
ACCESSION AM356079.1 GI:6860085
VERSION
KEYWORDS
SOURCE
ORGANISM

Bos taurus
Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovinae; Bos.
1 (bases 1 to 399)
Smith,T.P.L., Grosse,W.M., Freking,B.A., Roberts,A.J., Stone,R.T.,
Casas,E., Wray,J.E., White,J., Cho,J., Fahrenkrug,S.C., Bennett,
G.L., Heaton,M.P., Laegreid,W.M., Rohrer,G.A., Chitko-McKown,C.G.,
Perta,G., Holt,I., Karamycheva,S., Liang,F., Quackenbush,J. and
Keefe,J.W.
Sequence evaluation of four pooled-tissue normalized bovine CDNA
libraries and construction of a gene index for cattle
Genome Res. 11 (4), 626-630 (2001)
21180013

TITLE
JOURNAL
MEDLINE
COMMENT
Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 20
and -mismatch 12 options.
PCR Primers
FORWARD: AGGAACAGTATGACCAT
BACKWARD: GTTTCCAGTCACGACG
Plate: 17 row: P column: 9
Seq primer: ATTACGTGACACTATAG.
Location/Qualifiers
1..399
/organism="Bos taurus"
/db_xref="taxon:9913"

FEATURES

source

/clone_lib="MARC 2BOV"
 /tissue_type="pooled"
 /lab_host="DH10B"
 /note="Vector: PCMV SPORF6; Site_1: XbaI; Site_2: XhoI;
 library made from pooled tissue from testis, thymus,
 semitendinosus muscle, longissimus muscle, pancreas,
 adrenal, and endometrium."

BASE COUNT 100 a 126 c 117 g 56 t

Query Match 66.7% Score 67.4; DB 9; Length 399;
 Best Local Similarity 79.2%; Pred. No. 1.8e-07;
 Matches 80; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

OY 1 gggccaagccttgatgagacagcagaagtcggtgaggaaggagcaccagc 60
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 122 GGACCAAGGCTCTCAAGATGGAACCCAGAGATGTAGAGAAAGGAGCTGCCAGCC 181
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 OY 61 agggggaaggtcccgctccagcagcagctgctgagtcacc 101
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 182 TGGAGGAGGCTCCAGTCCAGGAGCTGCTGATCCACC 222

RESULT 2

AM427435 422 bp mRNA linear EST 25-APR-2001
 LOCUS 63185 MARC 3BOV Bos taurus cDNA 5', mRNA sequence.
 DEFINITION AM427435
 ACCESSION AM427435.1 GI:6955382
 VERSION EST.
 KEYWORDS

SOURCE
 ORGANISM

Bos taurus
 Eukaryota; Metazoa; Chordata; Cranialia; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Cetartiodactyla; Kuminantia; Pecora; Bovidae;
 Bovidae; Bovinae; Bos.
 1 (bases 1 to 422)
 Smith,T.P.L., Grosse,W.M., Freking,B.A., Roberts,A.J., Stone,R.T.,
 Casas,E., Wray,J.E., White,J., Cho,J., Fahrenkrug,S.C., Bennett,
 G.L., Heaton,M.P., Laegreid,W.W., Rohrer,G.A., Chiklo-Mckown,C.G.,
 Pettea,G., Holt,L., Karanycheva,S., Liang,F., Quackenbush,J. and
 Keefe,J.W.

Sequence evaluation of four pooled-tissue normalized bovine cDNA
 libraries and construction of a gene index for cattle
 Genome Res. 11 (4), 626-630 (2001)
 21180013
 CONTACT: Smith TPL
 USDA, ARS, US Meat Animal Research Center
 PO Box 166, Clay Center, NE 68933-0166, USA
 Tel: 402 762 4366
 Fax: 402 762 4390
 Email: smitht@mail.marc.usda.gov

Single pass sequencing. Bases called and trimmed with phred
 v0.980904.e. Vector identified by cross-match with the -minscore 20
 and -mismatch 12 options.
 PCR primers
 FORWARD: AGGAACAGCTATGACCAT
 BACKWARD: GTTTCACGATCAGCAGC
 Plate: 32 row: F column: 22
 Seq primer: ATTACGTGACACTATAG.
 Location/Qualifiers

FEATURES

source

/organism="Bos taurus"
 /db_xref="taxon:9913"
 /clone_lib="MARC 3BOV"
 /tissue_type="pooled"
 /lab_host="DH10B"
 /note="Vector: PCMV SPORF6; Site_1: XbaI; Site_2: XhoI;
 library made from pooled tissue from marrow, alveolar
 macrophage, ovary, fetal semitendinosus muscle, and fetal
 longissimus muscle."
 BASE COUNT 102 a 136 c 125 g 59 t

Query Match 66.7% Score 67.4; DB 9; Length 422;
 Best Local Similarity 79.2%; Pred. No. 1.8e-07;
 Matches 80; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

OY 1 gggccaagccttgatgagacagcagaagtcggtgaggaaggagcaccagc 60
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 170 GGACCAAGGCTCTCAAGATGGAACCCAGAGATGTAGAGAAAGGAGCTGCCAGCC 229
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 OY 61 agggggaaggtcccgctccagcagcagctgctgagtcacc 101
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 DB 230 TGGAGGAGGCTCCAGTCCAGGAGCTGCTGATCCACC 270

RESULT 3

BB628877 621 bp mRNA linear EST 31-AUG-2001
 LOCUS BB628877
 DEFINITION BB628877 RIKEN full-length enriched, 16 days neonate cerebellum Mus
 musculus cDNA clone 9630039L22 5', mRNA sequence.
 ACCESSION BB628877
 VERSION BB628877.1 GI:15399483
 KEYWORDS

SOURCE
 ORGANISM

Mus musculus
 Eukaryota; Metazoa; Chordata; Cranialia; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 621)
 Arakawa,T., Carninci,P., Fukuda,S., Furuno,M., Hanagaki,T., Hara,A.,
 Hiramoto,K., Hori,F., Ishii,Y., Ito,M., Kawai,J., Kono,H., Kouda,
 M., Koya,S., Matsuyama,T., Miyazaki,A., Nomura,K., Ohno,H., Okada,
 Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sano,H., Sasaki,
 D., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H.,
 Tagami,M., Tagawa,A., Takahashi,F., Takeda,Y., Tanaka,T., Toya,T.,
 Muramatsu,M. and Hayashizaki,Y.

RIKEN Mouse ESTs (Arakawa,T., et al. 2001)
 RIKEN Mouse ESTs (Arakawa,T., et al. 2001)
 Unpublished (2001)
 CONTACT: Yoshinide Hayashizaki
 Laboratory for Genome Exploration Research Group, RIKEN Genomic
 Sciences Center(GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suenryo-cho, Tsunumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel: 81-45-503-9222
 Fax: 81-45-503-9216
 Email: genome-res@gsr.riken.go.jp,
 URL: http://genome.gsc.riken.go.jp/
 Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K., Itoh,
 M., Kono,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.

Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new
 genes. Genome Res. 10 (10), 1617-1630 (2000)
 Wagi,K., Fujiwara,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E.,
 Watanabe,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsura,
 S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and
 Hayashizaki,Y.

RIKEN integrated sequence analysis (RISA) system--384-format
 sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)
 Kono,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara,
 Y. and Hayashizaki,Y.
 Computer-based methods for the mouse full-length cDNA
 encyclopedia: real-time sequence clustering for construction of a
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
 Yamanka,I., Kiyosawa,H., Kondo,S., Saito,T., Shinagawa,A., Aizawa,
 K., Fukuda,S., Hara,A., Itoh,M., Kawai,J., Shibata,K., Arakawa,T.,
 Ishii,Y. and Hayashizaki,Y.
 Mapping of 19032 mouse cDNAs on mouse chromosomes. J. Struct.
 Func. Genomics 2 pre, 172-186 (2001)
 Please visit our web site (http://genome.gsc.riken.go.jp) for
 further details.
 e mouse tissues.
 Location/Qualifiers

FEATURES

source

1. 621

```

/organism="Mus musculus"
/db_xref="taxon:10090"
/cclone="630039122"
/clone_1ib="RIKEN full-length enriched, 16 days neonate
cercobellum"
/tissue_type="cerebellum"
/dev_stage="16 days neonate"
/lab_host="DH10B"
/notes="Site_1: Salt; Site_2: BamHI. cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
primed with a primer 15'
GAGCAGAAGACCATCCAAAGACTCTTTTATTTTTTTTTTTTNN 3'. cDNA was
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. cDNA went through one round of normalization
to Rot = 20.0 and subtraction to Rot = 370.0. Second
strand cDNA was prepared with the primer adapter of
sequence 15' GAGCAGAAGATTCTCGAGTGTAATTAATTCACCCCCCCC
3}. cDNA was cleaved with xhoI and BamHI. Vector: a
modified pBluescript KS(+) after bulk excision from Lambda
FLC I."

BASE COUNT      143 a      183 g      101 t
ORIGIN

Query Match          66.7% Score 67.4; DB 9; Length 621;
Best Local Similarity 79.2%; Pred. No. 2e-07;
Matches 80; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

OY    1 ggagccaaagcttgatgacaaaggatcggtggagaaggagaccagagtc 60
      | ||| ||| ||||| ||||| || ||| ||| ||||| ||||| |||
Db    203 GAGTCAAGGCTTCAGATGACGACGACGAGGAGGCCGTAGAGAACACAGACCAGATT 262
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||
OY    61 aggggaagagtcgcccggtccagagccaagtgcgtgatccaac 101
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||
Db    263 tcggagaaaggtcccagttccagaccagactcgtccattcacc 303
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RESULT 4
BB629521         655 bp mRNA linear EST 31-AUG-2001
LOCUS           BB629521 RIKEN full-length enriched, adult male bone Mus musculus
DEFINITION     CDNA clone 9830138C07 5', mRNA sequence.
ACCESSION      BB629521
VERSION        BB629521.1 GI:15399646
KEYWORDS       EST.
SOURCE         house mouse.
ORGANISM       Mus musculus
REFERENCE      Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
                Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
                1 (bases 1 to 655)
AUTHORS       Arakawa,T., Carinici,P., Fukuda,S., Furuno,M., Hanagaki,T., Hara,A.,
                Hiramoto,K., Horii,F., Ishii,Y., Ito,M., Kawai,J., Konno,H., Kouda
                Okazaki,T., Okido,T., Saito,R., Sakai,C., Sakai,K., Sano,H., Sasaki
                ,D., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagami,M.,
                Tagawa,A., Takahashi,F., Takeda,Y., Tanaka,T., Toya,T., Muramatsu,M. and
                Hayashizaki,Y.
                RIKEN Mouse ESTs (Arakawa,T., et al. 2001)
                Unpublished (2001)
CONTACT       Yoshinhide Hayashizaki
                Laboratory for Genome Exploration Research Group, RIKEN Genomic
                Sciences Center(GSC), Yokohama Institute
                The Institute of Physical and Chemical Research (RIKEN)
                1-7-22 Suehiro-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan
                Tel.: 81-45-503-9222
                Fax: 81-45-503-9216
                Email: genome-res@sc.riken.go.jp,
                URL:http://genome.gsc.riken.go.jp/
```

Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Kono, H., Okazaki, Y., Muramatsu, M., and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. *Genome Res.* 10 (10), 1617-1630 (2000)

Wagti, K., Fujitake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Matsuki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsunura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kita, A., and Hayashizaki, Y.

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res.* 10 (11), 1757-1771 (2000)

Kono, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y., and Hayashizaki, Y.

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)

Yamanaka, I., Kiyosawa, H., Kondo, S., Saito, T., Shinagawa, A., Aizawa, K., Fukuda, S., Hata, A., Itoh, M., Kawai, J., Shibata, K., Araiawa, T., Ishii, Y., and Hayashizaki, Y.

Mapping of 19032 mouse cDNAs on mouse chromosomes. *J. Struct. Func. Genomics* 2 pre, L72-L86 (2001)

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

e mouse tissues.

FEATURES					
	source	location/Qualifiers			
		1..655	/organism="Mus musculus"		
		/db_xref="taxon:10090"			
		/clone="9830138C07"			
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		/dev_stage="adult"			
		/lab_host="DH10B"			
		/note="Site_1: SalI; Site_2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer 15'			
		GAGAGGAGAGAGATCCAAAGACCTCTTTTTCCTTTTTCVN 3'. cDNA was prepared by using trehalase thermo-activated reverse transcriptase and subsequently enriched for full-length b cap-trapper. cDNA went through one round of normalization to Rot - 10.0 and subtraction to Rot = 185.0. Second strand cDNA was prepared with the primer adapter of sequence 15' GAGAGAGATTCTTCGACTTAAATTAATCCCCCCCC 3'. cDNA was cleaved with XhoI and BamHI. Vector: a modified pBluescript KS(+) after bulk excision from Lambda			
		FLC 1."			
BASE COUNT	148 a	204 c	202 g	101 t	
ORIGIN					
Query Match	66.7%	Score 67.4:	DB 9:	Length 655:	
Best Local Similarity	79.2%:	Pred No. 2e-07:			
Matches 80;	Conservative 0;	Mismatches 21;	Indels 0;	Gaps 0	
OY 1	gggccaaagccttgatgacgaagcgagaagtcgltgaggaaaggagccaccagtc 60				
	I I				
Db 264	GAGTCAGCGCTTCACATGCAGCAGACGAGAGAGGCCCTGAGAGGAAGCAAGCACAGGT 323				
	I I				
OY 61	agggggaaaggtcccgcgttcaggccaagctgctgagtccacc 101				
	I I				
Db 324	TGGGAGAAGGTGCCAGTCACAGACCAGCTGTAATCCACC 364				
RESULT 5					
LOCUS	BB630381	685 bp	mRNA	linear	EST 26-OCT-2001
DEFINITION	BB630381 RIKEN full-length enriched, 6 days neonate skin Mus				

Query Match	54.1%: Score 54.6: DB 10: Length 548:
Best Local Similarity	77.6%: Pred. No. 0.00033:
Matches	66: Conservative 0: Mismatches 19: Indels 0: Gaps 0:
QY 17 alggacagagcagaatcgcgtlqgaagaagggagacaccagaagtcagagggaaggtccccg 76	
Dh 33 ATGCAAAAGCCACGAGGAGTGTAAACGAAAGGAGTGTCCAGGCGTGGAGGAGGTCCCA 92	
QY 77 gllccagagccagctgtgtagtcacc 101	
Dh 93 TTTCGAAGGCACTCTGTGAGTCCACC 117	

RESULT	7
A1664508	
LOCUS	
DEFINITION	A166A508 516 bp mRNA linear EST 10-MAY-1999
ACCESSION	U25005.Y1 Sugano mouse embryo mewa Mus musculus cDNA clone
VERSION	A166A508 IMAGE:1970001 5', mRNA sequence.
KEYWORDS	A166A508.1 GI:4768091
SOURCE	EST.
ORGANISM	house mouse. Mus musculus

REFERENCE AUTHORS

TITLE	The WashU-NCI Mouse EST Project 1999
JOURNAL	Unpublished (1999)
COMMENT	Other_ESTs: wk25b05.x1

CONTACT: Maria M/wasnu-MCI Mouse EST Project 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MCI:986741
Seq primer: custom primer used
High quality sequence stop: 484.

FEATURES

BASE COUNT	ORIGIN
113 a	<pre> 1..516 /organism="Mus musculus" /strain="C57BL" /db_xref="taxon:10090" /clone_image:1970001" /clone_11b="Sugano mouse embryo mewa" /dev_stage="embryo, 14 dpc" /lab_host="DH10B" /notice="Vector: pME18-FL3; Site_1: DraIII (CAGCTGTC); Site_2: DraII (CAGCATGTC); 1st strand cDNA was primed with an oligo(dT) primer [ATGTGGCCCTTTTTTTTTTTTTT]; double-stranded cDNA was ligated to a [TTGTGGCCCTACTG], digested and cloned into distinct DraIII adaptor sites of the pME18-FL3 vector (5' site CAGCTGTC, 3' site CAGCATGTC). XhoI should be used to isolate the cDNA insert. Size selection was performed to exclude fragments <1.5kb. Library constructed by Dr. Sumio Sugano (University of Tokyo Institute of Medical Science). Custom primers for sequencing: 5' end primer CTTCTGCTCTAAAGCTGGC and 3' end primer CGACTCTCAGCTCCAGCACA." </pre>
139 c	171 g 90 t 3 others

Query Match

53.38; Score 53.8; DB 9; Length 516;

	Best Local Similarity	77.2%;	Pred.	No. 0.00051;	
Matches 78:	Conservative 0:	Mismatches 22:	IndeIs 1:	Gaps 1:	
OY	1	gggcccaagccttgcgtgatgacaaaggacaagaatcgctgaaggaagggaccaccagtgc	60		
Dd	408	GAGTAAAGGCTTCACGGATGTGAAGAACAAGAGGCCCGTAGAGCAACAGACCACCCAGATT	467		
OY	61	aaggggaagtcctccgcgtlccaagcccagcttgcatattcaccc	101		
Dd	468	TGGGAGAGGGTGCCA - GTCCAACACACTCCTCGTAATCCCACC	507		

[illegible]

REFERENCE AUTHORS

TITLE	Sequence evaluation of four pooled-tissue normalized bovine cDNA libraries and construction of a gene index for cattle
JOURNAL	Genome Res. 11 (4), 626-630 (2001)
MEDLINE	21180013
COMMENT	Contact: Smith TPI

USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.960904.e. Vector identified by cross_match with the -mismatch
and -mismatch 12 options.
PCR primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCACAGTCACGACG
Plate: 57 row: H column: 11
Seq primer: ATTTAGCTGACACTTAG.

FEATURES
Source

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/organism="Bos taurus"
/db_xref="taxon:9913"
/clone_lib="MARC 3B0V"
/tissue_type="pooled"
/lab_host="DH10B"
/note="Vector: pCMV SPORT6; Site_1: XbaI; Site_2: XhoI;
library made from pooled tissue from marrow, alveolar
macrophage, ovary, fetal semitendinosus muscle, and fetal
longissimus muscle."
BASE COUNT
ORIGIN
91 a      147 c      139 g      67 t

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Query Match	52.18;	Score 52.6;	DB 10;	Length 444;
Best Local Similarity	81.38;	Pred. No. 0.00099;		
Matches 61; Conservative	0;	Mismatches 14;	Indels 0;	Gaps 0;

[illegible]

Db 61 GCTGCTGATGTCACC 75

RESULT 9
LOCUS B1754053/c
DEFINITION 603027607F1 NIH_MGC_114 Homo sapiens cDNA clone IMAGE:5198016 5', mRNA sequence.
ACCESSION B1754053
VERSION B1754053
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 922)
AUTHORS NIH-MGC <http://mhc.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
DNA Sequencing by: Inocyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Plate: LLM11495 row: n column: 01
High quality sequence stop: 830.
Location/Qualifiers
1..922
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/db_xref="taxon:9606"
/clone_image="5198016"
/clone_lib="NIH_MGC_114"
/lab_host="DH10B"
/note="Organ: brain; Vector: pCMV-SPORT6; Site: 1: NotI; Site: 2: EcoRV (destroyed); RNA source anonymous pool of 6 male brains, age range 23-27 yo. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.5 kb, insert size range 1-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 019. Note: this is a NIH_MGC Library."

BASE COUNT 216 a 268 c 284 g 154 t

Query Match 35.2%; Score 35.6; DB 10; Length 922;
Best Local Similarity 62.2%; Pred. NO. 23;
Matches 56; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

QY 4 ccaagcccttgatggaagaagcagtcggtggaaggaaggaagccaccagtcagg 63
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Db 915 CCACAGGTTTGGCCCGGATGGCTTCGAAATCCCGAGTGGAGTGGACCAACAGGGCCCT 856
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QY 64 gggaaagtcgcccgatccaggaagcagtcgctg 93
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Db 855 GGGAGTGGGCGCCGCTTCGCGCACCTGCTG 826
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RESULT 10
LOCUS BB759831/c
DEFINITION 406 bp mRNA linear EST 25-SEP-2001
clone G270121L23 3', mRNA sequence.
ACCESSION BB759831
VERSION BB759831
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 406)
AUTHORS Akimura, T., Atakawa, T., Carninci, P., Furuno, M., Hanagaki, T., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Imocani, K., Ishii, Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Nunase, R., Okazaki, Y., Okido, T., Salto, R., Sakai, C., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Tanaka, T., Tomaru, A., Toya, T., Watabiki, A., Yasunishi, A., Muramatsu, M., and Hayashizaki, Y.
TITLE RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)
JOURNAL Unpublished (2001)
COMMENT Contact: Yoshitake Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@sc.riken.go.jp,
URL:http://genome.gsc.riken.go.jp/,
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M., and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
wagi, K., Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watabiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A., and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system-384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y., and Hayashizaki, Y.
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.
e mouse tissues.
Location/Qualifiers
1..406
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone_image="G270121L23"
/clone_lib="RIKEN full-length enriched, melanocyte"
/cell_type="melanocyte"

BASE COUNT 77 a 130 c 98 g 101 t

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Best Local Similarity 58.9%; Pred. NO. 1.1e+02;
Matches 56; Conservative 0; Mismatches 39; Indels 0; Gaps 0;

QY 7 aagccttgatggaagaagcagtcggtggaaggaaggaagccaccagtcagg 66
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Db 257 AAGCCCTGAGTTCCCAAGCAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 198
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QY 67 aagtcgcccgatccaggaagcagtcgagtcacc 101
|||||

Db 197 CAGGAGCCCGAGCAGAGTGTCTCAAGTTCAAGC 163
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RESULT 11
LOCUS BB780864/c
DEFINITION 457 bp mRNA linear EST 15-NOV-2001
clone G430068010 3', mRNA sequence.
ACCESSION BB780864
VERSION BB780864
KEYWORDS EST.
SOURCE musculus cDNA clone G430068010 3', mRNA sequence.

ACCESSION VERSION KEYWORDS SOURCE ORGANISM	BB780864 BB780864.1 EST. house mouse. Mus musculus	GI:16941564
REFERENCE AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 457) Akihiro, T., Atakawa, T., Carninci, P., Furuno, M., Hanganaki, T.,	

TITLE	RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)
JOURNAL	Unpublished (2001)
COMMENT	Contact: Yoshihide Hayashizaki

Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suhei-to-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan
Tel.: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-resgsc.riken.go.jp,
URL: <http://genome.gsc.riken.go.jp/>
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh
, M., Kono, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. *Genome Res.* 10 (10), 1617-1630 (2000)
Wagii, K., Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
Wachiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura
, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kita, A. and
Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. *Genome Res.*
10 (11), 1757-1771 (2000)
Kono, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara
, Y. and Hayashizaki, Y.
Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)
Please visit our web site (<http://genome.gsc.riken.go.jp/>) for
further details.
e mouse tissues.

FEATURES	location/Qualifiers
source	1. .457

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/organism="Mus musculus"
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C1e-H3 cDNA"
/tissue_type="colon"
/cell_line="RCB-0549 C1e-H3"
BASE COUNT      93 a      145 c      106 g      113 t
ORIGIN

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Query Match	32.38;	Score 32.6;	DB 9;	Length 457;
Best Local Similarity	58.98;	Pred. No. 1.1e+02;		
Matches	56;	Conservative	0;	Mismatches 39;
			Indels	0;
			Gaps	0;

[illegible]

RESULT	12
LOCUS	B1765906/c
DEFINITION	B1765906 736 bp mRNA
ACCESSION	603046160F1 NIH_MGC_116 Homo sapiens cDNA clone IMAGE:5186367 5',
VERSION	B1765906
KEYWORDS	B1765906.1 GI:15757484
SOURCE	EST.
ORGANISM	human.
	Homo sapiens

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1 (bases 1 to 736)	NIH-MGC http://mgc.ncl.nih.gov/	National Institutes of Health, Mammalian Gene Collection (MGC)	Unpublished (1999)	Contact: Robert Strausberg, Ph.D.

Tissue: Cgagpds-remail.nln.gov
 Tissue Procurement: Life Technologies, Inc.
 cDNA Library Preparation: Life Technologies, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution by: MGC clone distribution
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 File: L1AM1465 row: 4 column: 16
 High quality sequence step: 736.

FEATURES

SOURCE

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/organism="Homo sapiens"
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/clone_1ID="NIH_MGC_116"
/_lab_host="DH10B"
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pCMV-SPORT6; Site_1: NotI; Site_2: EcoRV (destroyed); RNA
source anonymous pool of 3 colons, age 26 yo male, 49 yo
female, 71 yo male colon; 46 yo male kidney, and pool of 2
stomachs, 62 yo male and 70 yo female. Library is
oligo-dT primed and directionally cloned (EcoRV site is
destroyed upon cloning). Average insert size 1.4 kb,
insert size range 1-3 kb. Library is normalized and
enriched for full-length clones and was constructed by C.
Gruber (Invitrogen). Research Genetics tracking code
023. Note: this is a NIH_MGC Library."

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Query Match	32.38;	Score 32.6;	DB 10;	Length 736;
Best Local Similarity	60.98;	Pred. No. 1.2e+02;		
Matches 53;	Conservative 0;	Mismatches 34;	Indels 0;	Gaps 0

2 ggcacaagacctbtagtctgacacagtcagaaagtcgctgagggaggaaggagagccacaggta 6
 113 ggctaaagactccgtagagggggccagggccgacacggccgctcgagacacacaggggcc 54
 62 ggaggagagtctcccgctacagccagc 88
 53 agagacagaaagctcccgaggccacgctgc 27

RESULT	13
LOCUS	BG244994
DEFINITION	BG244994 151 bp mRNA linear EST 13-FEB-2001
	602358260F1 NCI_CCAP_Mam1 Mus musculus CDNA clone IMAGE:486719 5',
	mRNA sequence.

ORGANISM	SOURCE	KEYWORDS	VERSION	ACCESSION
Mus musculus	house mouse.	EST.	GI:127548099	BC244994.1
				BC244994

REFERENCE Mammalia: Eutheria: Rodentia: Sciurognathi: Muridae: Murinae: Mus.
 AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov

Tissue Procurement: Gilbert Smith, Ph.D.
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNLN at:
<http://image.llnl.gov>
 Plate: LLM10330 Row: h Column: 16
 High quality sequence stop: 151.

FEATURES
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 /strain="FVB/N"
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 /clone_image="NCI-CGAP_Mam1"
 /clone_id="NCI-CGAP_Mam1"
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 /lab_host="DH10B"
 /note="Organ: mammary; Vector: pCMV-SPORT6; Site_1: SalI;
 Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
 Library constructed by Life Technologies. Investigator
 providing samples: Gilbert Smith, NIH"

BASE COUNT 21 a 38 c 70 g 22 t
 ORIGIN

Query Match 32.1%; Score 32.4; DB 10; Length 151;
 Best Local Similarity 68.2%; Pred. No. 97;
 Matches 45; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

OY 14 gagatgacagcagaagtcggtggaggaaggaaccaccagtcagggaggaagtc 73
 Db 66 GAGGTGACCTCGCAGACGCCGTGGAGGCTGAGATCACCCGGAAGGGCGGACGACG 125
 OY 74 ccggtc 79
 Db 126 TGCCTC 131

RESULT 14
 BB860266 454 bp mRNA linear EST 26-NOV-2001
 LOCUS BB860266 RIKEN full-length enriched, pooled cell lines Mus musculus
 DEFINITION CDNA clone G430012004 5', mRNA sequence.
 ACCESSION BB860266
 VERSION BB860266.1 GI:17101720
 KEYWORDS EST.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 454)
 AUTHORS Akimura,T., Arikawa,T., Carninci,P., Furuno,M., Hanaoka,T.,
 Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,T., Imotani,K., Ishii,
 Y., Ito,M., Kawai,J., Kojima,Y., Kono,H., Kouda,M., Matsuyama,T.,
 Nakamura,M., Nishi,K., Nomura,K., Nunasaki,R., Okazaki,Y., Okido,T.,
 Saito,R., Sakai,C., Sakai,K., Sakazume,N., Sasaki,D., Sato,K.,
 Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa,
 A., Takahashi,F., Takaku-Akahira,S., Tanaka,T., Tomaru,A., Toya,T.,
 Watanabe,A., Yasunishi,A., Muramatsu,M. and Hayashizaki,Y.
 RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al.
 2001)
 JOURNAL Unpublished (2001)
 COMMENT Contact: Yoshihide Hayashizaki

Laboratory for Genome Exploration Research Group, RIKEN Genomic
 Sciences Center(GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel: 81-45-503-9222
 Fax: 81-45-503-9216

Email: genome-res@sc.riken.go.jp,
URL:http://genome.gsc.riken.go.jp/
 Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K., Itoh,
 M., Kono,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
 Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new
 genes. Genome Res. 10 (10), 1617-1630 (2000)
 wagi,K., Fujiwaka,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E.,
 Matsuhi,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsura
 S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and
 Hayashizaki,Y.

RIKEN integrated sequence analysis (RISA) system--384-format
 sequencing pipeline with 384 multichannel sequencer. Genome Res. 10 (11), 1757-1771 (2000)
 Kono,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara,
 Y. and Hayashizaki,Y.
 Computer-based methods for the mouse full-length cDNA
 encyclopedia: real-time sequence clustering for construction of a
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
 Please visit our web site (<http://genome.gsc.riken.go.jp>) for
 further details.
 e mouse tissues.

FEATURES
 Source

location/Qualifiers
 1..454
 /organism="Mus musculus"
 /db_xref="taxon:10090"
 /clone="G430012004"
 /clone_id="RIKEN full-length enriched, pooled cell lines"
 /note="pooled cell lines; (cell_line=RCB-0035 WEHI 164),
 (cell_line=RCB-2116 JC), (cell_line=RCB-0035 WEHI 164),
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 (cell_line=RCB-0559 K-1, F1), (cell_line=RCB-1283 B16
 melanoma), (cell_type=B cells), (cell_line=RCB-1702 WEHI 231
), (cell_type=Leydig cells), (cell_line=RCB-2065 M1TC-1),
 (cell_type=Nullipotent stem cell), (cell_line=RCB-2070 NE),
 (tissue_type=bladder, cell_line=RCB-0544 MBT-2),
 (tissue_type=bone marrow, cell_type=stroma cell,
 (tissue_type=bone marrow, cell_type=stroma cell,
 cell_line=CR-2028 SR-4987), (tissue_type=colon,
 cell_line=RCB-0549 C1e-H3), (tissue_type=kidney,
 cell_line=CCF-142 RAG), (tissue_type=submandibular gland,
 cell_line=CR-1734 SC4-9 clone 15), (strain=BA16/C,
 cell_type=B cells, cell_line=CR-1669 BCL1 clone 13, 20-3B3
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 BC3H1)"

BASE COUNT 121 a 123 c 133 g 77 t
 ORIGIN

Query Match 31.7%; Score 32; DB 9; Length 454;
 Best Local Similarity 60.2%; Pred. No. 1.6e+02;
 Matches 53; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

OY 6 aagacctgagatgacagcagaagtcggtggaggaaggaaccaccagtcaggg 65
 Db 357 ACACCATAGAGATGTCAAGGCAAGATCAGACAGAGGAGGATCCCTGACACGCA 416
 OY 66 gaaggtcccggtccagcagtcgctg 93
 Db 417 GAGCGTGAATCTTTCAGAGCAACACGACCTG 444

RESULT 15
 BF549656 479 bp mRNA linear EST 12-DEC-2000
 LOCUS BF549656
 DEFINITION UI-R-C2-ng-f-01-0-UI.r1 UI-R-C2 Rattus norvegicus cDNA clone
 ACCESSION BF549656
 UI-R-C2-ng-f-01-0-UI 5', mRNA sequence.

VERSION BP549656.1 GI:11659386
 KEYWORDS EST.
 SOURCE Norway rat.
 ORGANISM Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Rattus.

REFERENCE 1 (bases 1 to 479)
 AUTHORS Bonaldo,M.F., Lennon,G. and Soares,M.B.
 TITLE Normalization and subtraction: two approaches to facilitate gene
 discovery

JOURNAL Genome Res. 6 (9), 791-806 (1996)
 MEDLINE 97044477

COMMENT Contact: Soares, MB
 Program for Rat Gene Discovery and Mapping
 University of Iowa
 451 Eckstein Medical Research Building Iowa City, IA 52242, USA
 Tel: 319 335 8250
 Fax: 319 335 9565
 Email: msoares@blue.weeg.uiowa.edu

CDNA Library Preparation: M.B. Soares Lab Clone distribution:
 clones will be available through Research Genetics (www.resgen.com)
 This clone is also available through the I.M.A.G.E. Consortium at
 LBNL (info@image.lbnl.gov). IMAGE ID-1776352
 Seq primer: M13 Forward.

FEATURES
 source location/Qualifiers

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1..479
/organism="Rattus norvegicus"
/strain="Sprague-Dawley"
/db_xref="taxon:10116"
/clone="UI-R-C2-ng-f-0-UI"
/clone_1lb="UI-R-C2"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies)"
/note="Vector: pT73D-Pac (Pharmacia) with a modified
polylinker. Site_1: Not 1; Site_2: Eco RI; The UI-R-C2
library is a subtracted library derived from the UI-R-C1
library, which is a subtracted library derived from the
UI-R-C0 library. The UI-R-C0 library consisted of a
mixture of individually tagged normalized libraries
constructed from rat placenta, adult lung, brain, liver,
kidney, heart, spleen, ovary, muscle, 8, 12 and 18-day
embryo. The tag is a string of 3-5 nucleotides present
between the Not I site and the oligo-dT track which allows
identification of the library of origin of a clone within
the mixture. The subtracted library (UI-R-C2) was
constructed as follows: PCR amplified cDNA inserts from
UI-R-C1 clones from which 3' ESTs had been derived was
used as a driver in a hybridization with the UI-R-C1
library in the form of single-stranded circles. The
remaining single-stranded circles (subtracted library) was
purified by hydroxapatite column chromatography,
converted to double-stranded circles and electroporated
into DH10B bacteria (Life Technologies) to generate the
UI-R-C2 library. This procedure has been previously
described (Bonaldo, Lennon and Soares, Genome Research 6:
791-806, 1996)"
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BASE COUNT 103 a 163 c 146 g 65 t 2 others
 ORIGIN

Query Match 31.7%; Score 32; DB 10; Length 479;
 Best Local Similarity 62.5%; Pred. No. 1.6e+02;
 Matches 50; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

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OY 9 gccctgagatgacaagaagtcgtgaggaaggagccacgagtcaggagaa 68
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Db 48 GACTTGCGGCTGCACAAAGCAGACGCTGACACTGACATGAGCCAGCGGTGCC 107
   | | | | | | | | | | | | | | | | | | | | | | | | | |
OY 69 ggtcccggtccagccagc 88
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Db 108 AGAGCCCTCGGCAAGCGACG 127
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STATE: Massachusetts
COUNTRY: US
ZIP: 02173
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: IBM PC compatible
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/800,840
FILING DATE: 14-FEB-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/285,385
FILING DATE: 03-AUG-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/09905
FILING DATE: 03-AUG-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/776,859
FILING DATE: 03-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: Brook, David E.
REGISTRATION NUMBER: 22,592
REFERENCE/DOCKET NUMBER: MS797-01A2
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-861-6240
TELEFAX: 617-861-9540
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 268 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
US-08-800-840-1

Query Match      28.3% Score 28.6; DB 3; Length 268;
Best Local Similarity 55.6%; Pred.No. 2.6;
Matches 55; Conservative 0; Mismatches 44; Indels 0; Gaps 0.

Oy 1 gggcacaagccttgatgacgaaggcagaagtcgtgtaggaaggcgaccacccagtc 60
    ||||| |||| | ||| ||| | | | | | | | | | | | | | | | | |
Db 50 GAGCCACACCCTTAGGATGGCCAATCTACTCCAGAGCAGGAGCGGCAGACCGCAGGCC 109
    ||||| |||| | ||| ||| | | | | | | | | | | | | | | | | |

Oy 61 aggggaaagtgctccgcggtcccaagccagctgctcgagtcca 99
    ||||| |||| | ||| ||| | | | | | | | | | | | | | | | | |
Db 110 TGGGCATAAAAGTCAGGCGCAGACCATCTATTGCTTACA 148
    ||||| |||| | ||| ||| | | | | | | | | | | | | | | | | |

RESULT 3
US-08-776-859-1
Sequence 1, Application US/08776859
Patent No. 6090592
GENERAL INFORMATION:
APPLICANT: Mosaic Technologies, Inc.
APPLICANT: Adams, Christopher P.
TITLE OF INVENTION: Method and Apparatus for Performing
FILE REFERENCE: Mosaic
CURRENT APPLICATION NUMBER: US/08/776,859
CURRENT FILING DATE: 1997-05-29
EARLIER APPLICATION NUMBER: 08/285,385
NUMBER OF SEQ ID NOS: 4
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 1
LENGTH: 268
TYPE: DNA
ORGANISM: Homo sapiens
US-08-776-859-1
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Query Match      28.3%: Score 28.6; DB 3; Length 268;
Best Local Similarity 55.6%: Pred. No. 2.6;
Matches 55; Conservative 0; Mismatches 44; Indels 0; Caps 0

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Db      50  gagccacacccctaggtgttgcacatctaccctccagagacgaggaaggagccaggc 109

QY      61  agggggaaggtcccggttccagccagctgtcgtatca 99
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Db      110  tgggcataaaagtcagagcgacagccatctatgtctaca 148

RESULT      4
PCT-US95-09905-1
; Sequence 1, Application PC/TUS9509905
; GENERAL INFORMATION:
; APPLICANT:
; APPLICANT:
; APPLICANT:
; APPLICANT:
; TITLE OF INVENTION: METHOD AND APPARATUS FOR PERFORMING
; TITLE OF INVENTION: AMPLIFICATION OF NUCLEIC ACIDS ON SUPPORTS
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: WOLF, GREENFIELD & SACKS, P.C.
; STREET: 600 ATLANTIC AVENUE
; CITY: BOSTON
; STATE: MASSACHUSETTS
; COUNTRY: USA
; ZIP: 02210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/09905
; FILING DATE: FILED HEREMITH
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/285,385
; FILING DATE: 03-AUGUST-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: JANLUK, ANTHONY J.
; REGISTRATION NUMBER: 29,809
; REFERENCE/DOCKET NUMBER: B0833/7001WO
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-720-3500
; TELEFAX: 617-720-2441
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 268 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
PCT-US95-09905-1

Query Match      28.3%: Score 28.6; DB 5; Length 268;
Best Local Similarity 55.6%: Pred. No. 2.6;
Matches 55; Conservative 0; Mismatches 44; Indels 0; Caps 0;

QY      1  gggccaaagccttgatgacacaagcagaagtcggtggaaggaaggagccaccagtc 60
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Db      50  gagccacacccctaggtgttgcacatctaccctccagagacgaggaaggagccaggc 109

QY      61  agggggaaggtcccggttccagccagctgtcgtatca 99
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Db      110  tgggcataaaagtcagagcgacagccatctatgtctaca 148

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TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: Linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Homo sapiens
POSITION IN GENOME:
CHROMOSOME/SEGMENT: 11
PCT-US96-09430-7

Query Match 28.3% Score 28.6; DB 5; Length 6060;
Best Local Similarity 55.6% Pred. No. 4.7;
Matches 55; Conservative 0; Mismatches 44; Indels 0; Gaps 0;

Y 1 gggccaaagccttggagatggacaaagcgaagtcggtggaggaaggagccaccagtc 60
Db 2041 GAGCCACACCTTAGGCTTGGCCAAATCTACTCCAGGAGCAGGAGGAGGAGCCAGGCC 2100
Y 61 agggggaaggtcccggtccagggccagctgctgaatcca 99
Db 2101 TGGGCAATAAAGTCAGGCGCAGAGCATCTATTCCTTACA 2139

RESULT 11
US-08-289-653-2/c
Sequence 2, Application US/08289653
Patent No. 5543322
GENERAL INFORMATION:
APPLICANT: Kazuaki KITANO et al.
TITLE OF INVENTION: DNA AND ITS USE
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Wenderoth, Lind & Ponack
STREET: 805 Fifteenth Street, N.W., #700
CITY: Washington
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20005
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 500 kb
COMPUTER: IBM Compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/289, 653
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/07/887, 284
FILING DATE: May 22, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warren M. Cheek, Jr.
REGISTRATION NUMBER: 33,367
REFERENCE/DOCKET NUMBER:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-8850
TELEFAX:
TELEX:
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1140 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: Linear
MOLECULE TYPE: Genomic DNA
HYPOTHETICAL:
ANTI-SENSE:
FRAGMENT TYPE:
ORIGINAL SOURCE:
ORGANISM: Fusarium sp.

STRAIN: S-19-5 (IFO 8884)
INDIVIDUAL ISOLATE:
DEVELOPMENTAL STAGE:
HAPLOTYPE:
TISSUE TYPE:
CELL TYPE:
CELL LINE:
ORGANELLE:
IMMEDIATE SOURCE:
LIBRARY:
CLONE:
POSITION IN GENOME:
CHROMOSOME/SEGMENT:
MAP POSITION:
UNITS:
FEATURE:
NAME/KEY:
LOCATION:
IDENTIFICATION METHOD:
OTHER INFORMATION:
PUBLICATION INFORMATION:
AUTHORS:
TITLE:
JOURNAL:
VOLUME:
ISSUE:
PAGES:
DATE:
DOCUMENT NUMBER:
FILING DATE:
PUBLICATION DATE:
RELEVANT RESIDUES IN SEQ ID NO:

US-08-289-653-2

Query Match 26.5% Score 26.8; DB 1; Length 1140;
Best Local Similarity 57.0% Pred. No. 12;
Matches 49; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

Y 10 ccttgagatggacaagcagaagtcggtggaggaaggagccaccagtcagggggaag 69
Db 124 CTTGAGCTTGACATGCTACTTCCAGCAATGGCTGCGAACCAGCGCTCGAGGAGAG 65
Y 70 gtcccggtccagggccagctgctgaag 95
Db 64 GAGCGGCTTCAGGACCGGCGCCGCG 39

RESULT 12
US-08-289-653-1/c
Sequence 1, Application US/08289653
Patent No. 5543322
GENERAL INFORMATION:
APPLICANT: Kazuaki KITANO et al.
TITLE OF INVENTION: DNA AND ITS USE
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Wenderoth, Lind & Ponack
STREET: 805 Fifteenth Street, N.W., #700
CITY: Washington
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20005
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 500 kb
COMPUTER: IBM Compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/289, 653
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:


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: STREET: 345 California Street
: CITY: San Francisco
: STATE: California
: COUNTRY: USA
: ZIP: 94104-2675
:
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/164,292B
: FILING DATE: 09-DEC-1993
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: GRACEY, NANCY J.
: REGISTRATION NUMBER: 28,216
: REFERENCE/DOCKET NUMBER: 29310-20021.00
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (415) 677-7000
: TELEFAX: (415) 677-7522
: TELEX: 34-0154
: INFORMATION FOR SEQ ID NO: 17:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 5100 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 408..1331
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: US-08-164-292B-17
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: Best Local Similarity 61.4%; Pred. No. 16;
: Matches 43; Conservative 0; Mismatches 27; Indels 0; Gaps 0;
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: Db 1682 gcttgaagatggaagaagtcggtggaagaagggagccaccaggtcaagggggaa 68
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:
: QY 69 ggtcccggt 78
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: Db 1622 gctccctggt 1613
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: RESULT 15
: US-08-164-292B-19/C
: Sequence 19, Application US/08164292B
: Patent No. 5820868
:
: GENERAL INFORMATION:
: APPLICANT: MITTAL, SURESH K.
: APPLICANT: GRAHAM, FRANK L.
: APPLICANT: PREVEC, LUDVIK
: APPLICANT: BABIUK, LORNE A.
: TITLE OF INVENTION: RECOMBINANT PROTEIN PRODUCTION IN BOVINE
: TITLE OF INVENTION: ADENOVIRUS EXPRESSION VECTOR SYSTEM
: NUMBER OF SEQUENCES: 34
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: MORRISON & FOERSTER
: STREET: 345 California Street
: CITY: San Francisco
: STATE: California
: COUNTRY: USA
: ZIP: 94104-2675
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
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: APPLICATION NUMBER: US/08/164,292B
: FILING DATE: 09-DEC-1993
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: GRACEY, NANCY J.
: REGISTRATION NUMBER: 28,216
: REFERENCE/DOCKET NUMBER: 29310-20021.00
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (415) 677-7000
: TELEFAX: (415) 677-7522
: TELEX: 34-0154
: INFORMATION FOR SEQ ID NO: 19:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 5100 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 529..954
:
: US-08-164-292B-19
:
: Query Match          26.5%; Score 26.8; DB 1; Length 5100;
: Best Local Similarity 61.4%; Pred. No. 16;
: Matches 43; Conservative 0; Mismatches 27; Indels 0; Gaps 0;
:
: QY 9 gcttgaagatggaagaagtcggtggaagaagggagccaccaggtcaagggggaa 68
:    | | | | | | | | | | | | | | | | | | | | | | | | | | | |
: Db 1682 gcttgaagatggaagaagtcggtggaagaagggagccaccaggtcaagggggaa 68
:    | | | | | | | | | | | | | | | | | | | | | | | | | | | |
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: QY 69 ggtcccggt 78
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: Db 1622 gctccctggt 1613
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Search completed: October 3, 2002, 16:22:14
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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OW nucleic - nucleic search, using sw model

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(without alignments)

288,886 Million cell updates/sec

Title: US-09-826-581-5_COPY_500_600

Perfect score: 101
Sequence: 1 cgtcccccgcagggcccatl.....atgcagagacacacctgcta 101

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenBml: *
1: gb_ba: *
2: gb_hlg: *
3: gb_ln: *
4: gb_cm: *
5: gb_ov: *
6: gb_pat: *
7: gb_ph: *
8: gb_pl: *
9: gb_pr: *
10: gb_ro: *
11: gb_sts: *
12: gb_sy: *
13: gb_un: *
14: gb_vl: *
15: em_ba: *
16: em_fun: *
17: em_hum: *
18: em_ln: *
19: em_mu: *
20: em_om: *
21: em_or: *
22: em_ov: *
23: em_pat: *
24: em_ph: *
25: em_pl: *
26: em_ro: *
27: em_sts: *
28: em_un: *
29: em_vl: *
30: em_htg_hum: *
31: em_htg_inv: *
32: em_htg_other: *
33: em_htgo_inv: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
------------	-------	-------------	--------	-------	-------------

1	101	100.0	989	6	AX281579	AX281579 Sequence
2	101	100.0	1647	6	AX281582	AX281582 Sequence
3	101	100.0	2109	6	AX099776	AX099776 Sequence
4	101	100.0	2115	6	AX099802	AX099802 Sequence
5	101	100.0	2115	6	AF214519	AF214519 Homo sapi
6	101	100.0	152129	2	AC027416	AC027416 Homo sapi
7	101	100.0	196554	2	AC073128	AC073128 Homo sapi
8	97.8	96.8	2290	9	HS249977	HS249977 Homo sapi
9	89.6	88.7	206854	9	AC009974	AC009974 Homo sapi
10	72.4	71.7	1867	6	AX099774	AX099774 Sequence
11	72.4	71.7	1873	6	AF214520	AF214520 Sus scrofa
12	72.4	71.7	1873	6	AX099800	AX099800 Sequence
13	72.4	71.7	2022	6	AX099804	AX099804 Sequence
14	72.4	71.7	5888	4	AF214521	AF214521 Sus scrofa
15	72.4	71.7	227724	2	AF336381	AF336381 Mus muscu
16	33.4	33.1	3503	9	HS181675	HS181675 Homo sapien
17	33.4	33.1	81811	2	AC063981	AC063981 Homo sapi
18	33.4	33.1	82237	9	AC003677	AC003677 Human pl
19	33.4	33.1	108123	9	AC034206	AC034206 Homo sapi
20	33.4	33.1	159420	2	AC009017	AC009017 Homo sapi
21	32	31.7	790	33	AC076924	AC076924 Giardia
22	32	31.7	975	33	AC078451	AC078451 Giardia
23	32	31.7	988	33	AC028281	AC028281 Giardia
24	32	31.7	1927	3	AF071896	AF071896 Giardia
25	31.2	30.9	17570	1	AE007063	AE007063 Mycobacte
26	31.2	30.9	39430	1	MTCV49	MTCV49 Mycobacte
27	30.4	30.1	736	33	AC053985	AC053985 Giardia
28	30.4	30.1	804	33	AC053984	AC053984 Giardia
29	30.4	30.1	967	33	AC089076	AC089076 Giardia
30	30.4	30.1	1283	33	AF050755	AF050755 Giardia
31	30.4	30.1	1306	3	AF050754	AF050754 Giardia
32	30.4	30.1	3989	3	AF071897	AF071897 Giardia
33	30.4	30.1	5791	1	AE117694	AE117694 Rhizobium
34	30.2	29.9	11085	1	AE004450	AE004450 Pseudomon
35	29.6	29.3	3603	1	AF035395	AF035395 Pseudomon
36	29.6	29.3	3810	3	AC091120	AC091120 Leishman
37	29.4	29.1	850	33	AC042098	AC042098 Giardia
38	29.4	29.1	214609	2	AC024651	AC024651 Homo sapi
39	29.2	28.9	56425	7	AF165214	AF165214 Bacteriop
40	29	28.7	1575	3	AY069691	AY069691 Drosophil
41	29	28.7	69061	2	AC012986	AC012986 Drosophil
42	29	28.7	158628	2	AC020653	AC020653 Homo sapi
43	29	28.7	168469	3	AC007886	AC007886 Drosophil
44	29	28.7	173729	2	AC012236	AC012236 Homo sapi
45	29	28.7	228448	3	AE003772	AE003772 Drosophil

ALIGNMENTS

RESULT 1	AX281579	989 bp	DNA	linear	PAT 02-NOV-2001
LOCUS	AX281579	Sequence 2 from Patent WO0177305.			
DEFINITION	AX281579				
ACCESSION	AX281579.1	GI:16608830			
VERSION					
KEYWORDS	human.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Andersson, L., Luthman, H. and Marklund, S.				
AUTHORS	1 (sites)				
TITLE	Patent: WO 0177305-A 2 18-OCT-2001;				
JOURNAL	Arexis AB (SE)				
FEATURES	Location/Qualifiers				
source	1..989				
BASE COUNT	229 a 306 c 286 g 168 t				
ORIGIN	/organism="Homo sapiens"				
	/db_xref="taxon:9606"				

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Best Local Similarity 100.0%; Pred. No. 1.5e-17;
Matches 101: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 ctgtcccgagggcccaattcccaagctggctgagatgacgaactgcggaaccggc 60
793 CTGTCCCGAGGCCCAATTTCCCAAGCTGGCTGCGATGACGAACCTCGGAACCCGGC 852

Qy 61 gccagatctacatgcgtctcatatgcagagacacactgcta 101
683 GCCAGATCTACATGCGCTTCATGCAGACACACTCTCTA 893

RESULT 2
LOCUS AX281582 1647 bp DNA linear PAT 02-NOV-2001
DEFINITION Sequence 5 from Patent WO0177305.
ACCESSION AX281582
VERSION AX281582.1 GI:16608833
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
AUTHORS Andersson, L., Luthman, H. and Marklund, S.
TITLE Variants of the human amp-activated protein kinase gamma 3 subunit
JOURNAL Patent: WO 0177305-A 5 18-OCR-2001;
Axeis AB (SE)

FEATURES
source Location/Qualifiers
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/organism="Homo sapiens"
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20..1489
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/db_xref="GI:16608834"
/translation="MEPGLLEHALRTPSMSSLOGSEHQMSEFLSENSSMSPAYTS
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LSPOAFEPKLGMDELARKGAOIYMRPMENHCYDMATSSKLVTFDTMLERKAFPA
LVANGVRAAPLINDSKOSFVCHLTITDFILVLRKYRSPLVQIYELEQHKIETWREIY
LOGCFKPLVISISNDSELEFVAVTLIKNRHRLPVDPSGVNVLHILTHRLKFLHIF
GSLPRPSFLYRTIDJIGTFRDIAVLETAPIILALDIFDRLVSALPVNCEGOV
VGLYSRFDIHLAAOQTYNHLDMSEALRORTLCLEGLVSCOPHESLGEVIDRIARE
QVRLVLVDETQHLGVVSLDILQALVSPAGIDALGA"

BASE COUNT 346 a 502 c 462 g 337 t

ORIGIN

Query Match 100.0% Score 101: DB 6: Length 1647;
Best Local Similarity 100.0%; Pred. No. 1.4e-17;
Matches 101: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ctgtcccgagggcccaattcccaagctggctgagatgacgaactgcggaaccggc 60
500 CTGTCCCGAGGCCCAATTTCCCAAGCTGGCTGCGATGACGAACCTCGGAACCCGGC 559

Db 61 gccagatctacatgcgtctcatatgcagagacacactgcta 101
560 GCCAGATCTACATGCGCTTCATGCAGACACACTCTCTA 600

RESULT 3
LOCUS AX099776 2109 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 3 from Patent WO0120003.
ACCESSION AX099776
VERSION AX099776.1 GI:13538810
KEYWORDS
SOURCE human.

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
AUTHORS Andersson, L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Kogel-Gallard, C., Iannuccelli, N., Gellin, J., le Roy, P. and
Charidon, P.
TITLE Variants of the gamma chain of amp, dna sequences encoding the
same, and uses thereof
JOURNAL Patent: WO 0120003-A 3 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Lelf (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)

FEATURES
source Location/Qualifiers
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/organism="Homo sapiens"
/db_xref="taxon:9606"
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YRTIDJIGTFRDIAVLETAPIILALDIFDRLVSALPVNCEGOVGLYSRDI
HLAAOQTYNHLDMSEALRORTLCLEGLVSCOPHESLGEVIDRIAREQVRLVLYDE
TQHLGVVSLDILQALVSPAGIDALGA"

BASE COUNT 458 a 621 c 560 g 470 t

ORIGIN

Query Match 100.0% Score 101: DB 6: Length 2109;
Best Local Similarity 100.0%; Pred. No. 1.4e-17;
Matches 101: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ctgtcccgagggcccaattcccaagctggctgagatgacgaactgcggaaccggc 60
400 CTGTCCCGAGGCCCAATTTCCCAAGCTGGCTGCGATGACGAACCTCGGAACCCGGC 459

Db 61 gccagatctacatgcgtctcatatgcagagacacactgcta 101
460 GCCAGATCTACATGCGCTTCATGCAGACACACTCTCTA 500

RESULT 4
LOCUS AX099802 2115 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 29 from Patent WO0120003.
ACCESSION AX099802
VERSION AX099802.1 GI:13538836
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
AUTHORS Andersson, L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Kogel-Gallard, C., Iannuccelli, N., Gellin, J., le Roy, P. and
Charidon, P.
TITLE Variants of the gamma chain of amp, dna sequences encoding the
same, and uses thereof
JOURNAL Patent: WO 0120003-A 29 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Lelf (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)

FEATURES
source Location/Qualifiers
1..2115
/organism="Homo sapiens"
/db_xref="taxon:9606"
1..1395
/note="unnamed protein product"
/codon_start=1
/protein_id="CAC35801.1"
/db_xref="GI:13538837"

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: MIBR

```
Web site: http://www.seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
-----
Project Information
Center project name: 17458
Center clone name: 504_C_11
-----
Summary Statistics
Sequencing vector: M13; M77815: 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 135376 bases at least Q40
Consensus quality: 143264 bases at least Q30
Consensus quality: 146503 bases at least Q20
Insert size: 161000; agarose-fp
Insert size: 149029; sum-of-contigs
Quality coverage: 3.1 in Q20 bases; agarose-fp
Quality coverage: 3.3 in Q20 bases; sum-of-contigs
-----
NOTE: This is a 'working draft' sequence. It currently
* consists of 32 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1
* 1006 1105: contig of 1005 bp in length
* 1106 2402: contig of 1297 bp in length
* 2403 2502: gap of 100 bp
* 2503 3823: contig of 1321 bp in length
* 3824 3923: gap of 100 bp
* 3924 5020: contig of 1097 bp in length
* 5021 5120: gap of 100 bp
* 5121 6161: contig of 1041 bp in length
* 6162 6261: gap of 100 bp
* 6262 7547: contig of 1286 bp in length
* 7548 7647: gap of 100 bp
* 7648 9983: contig of 2336 bp in length
* 9984 10083: gap of 100 bp
* 10084 12556: contig of 2473 bp in length
* 12557 12656: gap of 100 bp
* 12657 15043: contig of 2387 bp in length
* 15044 15143: gap of 100 bp
* 15144 17123: contig of 1980 bp in length
* 17124 17223: gap of 100 bp
* 17224 19466: contig of 2243 bp in length
* 19467 19566: gap of 100 bp
* 19567 21928: contig of 2362 bp in length
* 21929 22028: gap of 100 bp
* 22029 24319: contig of 2291 bp in length
* 24320 24419: gap of 100 bp
* 24420 27059: contig of 2640 bp in length
* 27060 27159: gap of 100 bp
* 27160 30170: contig of 3011 bp in length
* 30171 30270: gap of 100 bp
* 30271 33968: contig of 3698 bp in length
* 33969 34068: gap of 100 bp
* 34069 38179: contig of 4111 bp in length
* 38180 38279: gap of 100 bp
* 38280 42366: contig of 4087 bp in length
* 42367 42466: gap of 100 bp
* 42467 46365: contig of 3899 bp in length
* 46366 46465: gap of 100 bp
* 46466 51285: contig of 4820 bp in length
* 51286 51385: gap of 100 bp
* 51386 55871: contig of 4486 bp in length
* 55872 55971: gap of 100 bp
* 55972 60595: contig of 4624 bp in length
* 60596 60695: gap of 100 bp
* 60696 66595: contig of 5900 bp in length
* 66596 66695: gap of 100 bp
* 66696 73218: contig of 6523 bp in length
* 73219 73318: gap of 100 bp
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FEATURES
Source
* 73319 77115: contig of 3797 bp in length
* 77116 77215: gap of 100 bp
* 77216 85022: contig of 7807 bp in length
* 85023 85122: gap of 100 bp
* 85123 93314: contig of 8192 bp in length
* 93315 93414: gap of 100 bp
* 93415 101193: contig of 7779 bp in length
* 101194 101293: gap of 100 bp
* 101294 113090: contig of 11797 bp in length
* 113091 113190: gap of 100 bp
* 113191 123496: contig of 10306 bp in length
* 123497 123596: gap of 100 bp
* 123597 137837: contig of 14241 bp in length
* 137838 137937: gap of 100 bp
* 137938 152129: contig of 14192 bp in length.
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/clone="RP11-504G11"
/clone_11b="RPC1-11 Human Male BAC"
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1106..2402
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2503..3823
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5121..6161
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6262..7547
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7648..9983
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12657..15043
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17224..19466
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22029..24319
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27160..30170
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Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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db 38135  ctgtcccccagggcccatcttcccaagctggagtgacgaactgcggaaccgcgc 38076
         |||||||

QY      61      gccagatcatatcgctcatcagagagccacticta 101
         |||||||
db 38075  gccagatctacatgccttccatgcacagcacaccttcta 38035
         |||||||

RESULT 7
AC073128/c      196554 bp      DNA      linear      HTG 21-FEB-2001
LOCUS      Homo sapiens chromosome 2 clone RP11-64705, WORKING DRAFT SEQUENCE,
DEFINITION      17 unordered pieces.
ACCESSION      AC073128
VERSION      AC073128.3 GI:13027579
KEYWORDS      HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULTOP.
SOURCE      human.
ORGANISM      Homo sapiens
               Eukaryota; Metazoa; Chordata; Craniala; Vertebrata; Euteleostomi;
               Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE      1 (bases 1 to 196554)
AUTHORS      Waterston,R.H.
TITLE      The sequence of Homo sapiens clone
JOURNAL      Unpublished
AUTHORS      2 (bases 1 to 196554)
TITLE      Waterston,R.H.
JOURNAL      Submitted (08-JUN-2000) Genome Sequencing Center, Washington
               University School of Medicine, 4444 Forest Park Parkway, St. Louis,
               MO 63108, USA
COMMENT      On Feb 21, 2001 this sequence version replaced gi:8469048.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
Project Information -----
Center project name: H.NH0647005
Summary Statistics -----
Sequencing vector: MJ3: 98%
Sequencing vector: plasmid: 0%
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 187795 bases at least Q40
Consensus quality: 190513 bases at least Q30
Consensus quality: 192099 bases at least Q20
Insert size: 200000; agarose-fp
Insert size: 194954; sum-of-contigs
Quality coverage: 5.58 in Q20 bases; agarose-fp
Quality coverage: 5.67 in Q20 bases; sum-of-contigs
----- NOTE -----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 17 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will

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```

* be preserved.
1 1157: contig of 1157 bp in length
1158 1257: gap of unknown length
1258 3600: contig of 2343 bp in length
3601 3700: gap of unknown length
3701 3701: contig of 1403 bp in length
5104 5203: gap of unknown length
5204 8524: contig of 3321 bp in length
8525 8625: gap of unknown length
8626 11856: contig of 3232 bp in length
11857 11956: gap of unknown length
11957 15783: contig of 3827 bp in length
15784 15883: gap of unknown length
15884 21906: contig of 6023 bp in length
21907 22006: gap of unknown length
22007 28887: contig of 6881 bp in length
28888 28987: gap of unknown length
28988 35255: contig of 6268 bp in length
35256 35355: gap of unknown length
35356 44642: contig of 9287 bp in length
44643 44742: gap of unknown length
44743 58275: contig of 13533 bp in length
58276 58375: gap of unknown length
58376 73816: contig of 15441 bp in length
73817 73916: gap of unknown length
73917 92140: contig of 18224 bp in length
92141 92240: gap of unknown length
92241 113337: contig of 21097 bp in length
113338 113437: gap of unknown length
113438 130325: contig of 16888 bp in length
130326 130425: gap of unknown length
130426 149287: contig of 18862 bp in length
149288 149388: gap of unknown length
149389 196554: contig of 47167 bp in length.
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1. 196554
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/db_xref="taxon:9606"
/chromosome="2"
/clone="RP11-64705"
1. 1157
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1258. 3600
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3701. 5103
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clone_end:77
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5204. 8524
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8625. 11856
/note="assembly_name:Contig21"
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15884. 21906
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22007. 28887
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28988. 35255
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58376. 73816
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73917. 92140
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130426. 149287

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Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 62634 cttgtcccccagcccccatttcccaagctggctggatgacgaactcgcgaacccgcg 62575
Oy 61 gccagatctacatgccttctatgcaagacacactgcta 101
Db 62574 gccacatctacatgccttctatgcaagacacactgcta 62534

RESULT 8
LOCUS      HSA249977      2290 bp      mRNA      linear      PRI 07-APR-2000
DEFINITION      Homo sapiens mRNA for AMP-activated protein kinase gamma 3 subunit
ACCESSION      AJ249977
VERSION      AJ249977.1 GI:6688200
KEYWORDS      AMP-activated protein kinase; AMPK gamma 3 gene; gamma 3 subunit.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 2290)
Cheung, P.C., Salt, I.P., Davies, S.P., Hardie, D.G. and Carling, D.
Characterization of AMP-activated protein kinase gamma-subunit
isoforms and their role in AMP binding
Biochem. J. 346 Pt 3, 659-669 (2000)
2016409
2 (bases 1 to 2290)
Carling, D.
Direct Submission
Submitted (12-OCT-1999) Carling D., Cellular Stress Group, MRC
Clinical Sciences Centre, Hammersmith Hospital, DuCane Road,
London, W12 0NN, UNITED KINGDOM
Location/Qualifiers
1..2290
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/gene="AMPK gamma 3"
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/function="AMP-activated protein kinase regulatory
subunit"
/codon_start=1
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/product="AMP-activated protein kinase gamma 3 subunit"
/protein_id="CAB65117.1"
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/translation="MEPGLHALRRTPSWSLGSEHQHSMPLFEOENSSWSPAYTS
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LSQAPFPKLGMDLRRKPGAOIYMRPIETHTCYDMATSSKLVITDMLETKKAFEA
LVANGVRAAPLMDSKOSPFYCMITPTFTLVLRHYRSPVLVYELQKTEWREIY
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GSLRPSPLKRTIQQDGLGTFNDLAVLELTAPILTALDIFVDRVSALPVNECCOV
VGLSRFDVTHLAQDQVYVNHLDMSVGEALRKRLCLEGVISCPHSLGVEIDRLARE
QVRLVAVDEFQHLGVSLDILQALVISPACIDPSGPEKI"
BASE COUNT      501 a 674 c 617 g 498 t
ORIGIN

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Query Match
Best Local Similarity 96.8%; Score 97.8; DB 9; Length 2290;
Matches 99; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 502 cttgtcccccagcccccatttcccaagctggctggatgacgaactcgcgaacccgcg 561
Oy 61 gccagatctacatgccttctatgcaagacacactgcta 101
Db 562 gccacatctacatgccttctatgcaagacacactgcta 602

RESULT 9
LOCUS      AC009974/c      206854 bp      DNA      linear      PRI 09-JAN-2002
DEFINITION      Homo sapiens BAC clone RP11-459119 from 2, complete sequence.
ACCESSION      AC009974
VERSION      AC009974.9 GI:16799058
KEYWORDS      HTG.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 206854)
Sulston, J.E. and Waterston, R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
99063792
2 (bases 1 to 206854)
Harris, A. and Cotton, M.
The sequence of Homo sapiens BAC clone RP11-459119
Unpublished (2001)
3 (bases 1 to 206854)
Waterston, R.H.
Direct Submission
Submitted (08-SEP-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 206854)
Waterston, R.H.
Direct Submission
Submitted (08-NOV-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 206854)
Waterston, R.H.
Direct Submission
Submitted (03-JAN-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
6 (bases 1 to 206854)
Waterston, R.
Direct Submission
Submitted (09-JAN-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Nov 8, 2001 this sequence version replaced g1:13431203.
-----
Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: saplens@wustl.wustl.edu
-----
Summary Statistics
Center project name: H_NH045119

```

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:

all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Moon, P.Y., Zhao, B., Frengen, E., Tateno, M., Calanese, J.J., and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://daccpac.med.buffalo.edu>)

VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-1077K22; the clone sequenced to the right is RP11-64705. Actual start of this clone is at base position 1 of RP11-459119; actual end is at base position 206854 of RP11-459119.

Data from AC079810 and AC073128 was used to finish this clone. AC009974. Polymorphisms have been identified between AC073128 and AC009974. A single plasmid region exists between 38812-38903. An unresolved tandem in the HERY SVA exists between 184390-185163. PCR suggests that approximately 1700 bps are missing.

FEATURES

source

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misc_feature 967..1071

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Search completed: October 3, 2002, 14:49:54
Job time: 12114 sec

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Query Match      71.7%: Score 72.4; DB 4; Length 5888;
Best Local Similarity 83.7%: Pred. No. 7.6e-10;
Matches 82: Conservative 0; Mismatches 16; Indels 0; Gaps 0;

QY 4 lccccgagggcccaattcccaagctgggctggatgacgaactgcggaaccggcgcc 63
      ||||| ||| | | ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 769 TCCCCAGAGGCTGCTGTACCCAGGCTGGCTGGATGATGCTGCAGAGCCGGGGCCC 828
QY 64 cagatctacatgcgctcatalgcagagacacactgcta 101
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 829 CAGGCTACATGCACCTTCATGCAGAGACACACCTGCTA 866

RESULT 15
AF336381/c 227724 bp DNA linear HTG 02-APR-2001
LOCUS Mus musculus chromosome 1 clone PAC510; PAC457, *** SEQUENCING IN
DEFINITION
ACCESSION AF336381
VERSION AF336381.1 GI:13507298
KEYWORDS HTG: HTGS_PHASE1.
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi:
AUTHORS Mammalia: Eutheria: Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE 1 (bases 1 to 227724)
JOURNAL Rump, A., Hayes, C., Brown, S.D.M. and Rosenthal, A.
REFERENCE Mouse chromosome 1 genomic sequence
AUTHORS Unpublished
JOURNAL 2 (bases 1 to 227724)
Rump, A.
COMMENT Direct Submission
Submitted (17-JAN-2001) Genome Analysis Institute of Molecular
Biotechnology, Beutenbergstr. 11, Jena 07745, Germany
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
      1 17869: contig of 17869 bp in length
      * 17870 17969: gap of unknown length
      * 17970 32746: contig of 14777 bp in length
      * 32747 32846: gap of unknown length
      * 32847 227724: contig of 194878 bp in length.
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ORIGIN
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Best Local Similarity 83.7%: Pred. No. 5.8e-10;
Matches 82: Conservative 0; Mismatches 16; Indels 0; Gaps 0;

QY 4 lccccgagggcccaattcccaagctgggctggatgacgaactgcggaaccggcgcc 63
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Db 8211 TCCCCAGAGGCCCCACTCTGCTGAGCTGGATGATGCAACTTCAGAACCCCGGAGCC 8152

QY 64 cagatctacatgcgctcatalgcagagacacactgcta 101
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PN XX
XX MO200177305-A2.
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XX 18-OCT-2001.
XX
XX 06-APR-2001; 2001WO-SE00765.
XX
XX 07-APR-2000; 2000US-195665P.
XX
XX (AREX-) AREXIS AB.
XX
XX Andersson L, Luthman H, Marklund S;
XX
XX MPI; 2001-657170/75.
XX
XX
XX New variants of human AMP-activated protein kinase gamma3 subunit
XX associated with a metabolic disease e.g. diabetes or obesity and method
XX for determining a risk estimate of diseases in subject by detecting the
XX variant -
XX
XX Example 1; Fig 2; 25pp; English.
XX
XX The sequences given in AAH43681-84 represents genomic fragments
XX encoding the human AMP-activated protein kinase gamma 3 subunit
XX (PRKAG3). Detecting the presence of the PRKAG3 DNA, or a variant,
XX is useful in determining a risk estimate of a metabolic disease,
XX such as diabetes or obesity, in a subject. The variation may occur
XX in exons 3, 4 or 10. In exon 3 variation may be a substitution of
XX a G for a C at nucleotide 320, resulting in the amino acid
XX substitution P71A; in exon 4 variation may be a substitution of a
XX T for a C at nucleotide 550; and in exon 10 variation may be a
XX substitution of a T for a C at nucleotide 1037, resulting in the
XX amino acid substitution R340W. There may also be nucleotide variation
XX in intron 6.
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XX Sequence 989 BP; 229 A; 306 C; 286 G; 168 T; 0 other;
XX
XX
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XX Best Local Similarity 100.0%; Pred. No. 9.8e-23;
XX Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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XX 1 ctgtcccgagagcccaattcccaagctggctggatgacgaactcggaaaccggc 60
XX ||||||||||||||||||||||||||||||||||||||||||||||||||||
XX Db 793 ctgtcccgagagcccaattcccaagctggctggatgacgaactcggaaaccggc 852
XX
XX 61 gcccgatctacatgcgtctcatgcagagcacacctgcta 101
XX ||||||||||||||||||||||||||||||||||||||||||||||||
XX Db 853 gcccgatctacatgcgtctcatgcagagcacacctgcta 893
XX
XX
XX RESULT 2
XX AAH43685
XX ID AAH43685 standard; cDNA; 1647 BP.
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XX AAH43685;
XX
XX 21-JAN-2002 (first entry)
XX
XX PRKAG3 cDNA.
XX
XX Human; AMP-activated protein kinase gamma 3 subunit; PRKAG3; variant;
XX metabolic disease; diabetes; obesity; substitution; ss.
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XX Homo sapiens.
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XX CDS 20..1489
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XX FT 230
XX FT /*tag= b
XX variation
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FT FT /note= "Causes P71A"
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FT FT /note= "Silent variation"
FT variation
FT 1037
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FT FT /label= "C1037T"
FT FT /note= "Causes R340W"
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XX MO200177305-A2.
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XX 18-OCT-2001.
XX
XX
XX 06-APR-2001; 2001WO-SE00765.
XX
XX 07-APR-2000; 2000US-195665P.
XX
XX (AREX-) AREXIS AB.
XX
XX Andersson L, Luthman H, Marklund S;
XX
XX MPI; 2001-657170/75.
XX
XX P-PSDB; QDB47679.
XX
XX
XX New variants of human AMP-activated protein kinase gamma3 subunit
XX associated with a metabolic disease e.g. diabetes or obesity and method
XX for determining a risk estimate of diseases in subject by detecting the
XX variant -
XX
XX Disclosure; Fig 5; 25pp; English.
XX
XX This sequence represents the full length cDNA encoding the human
XX AMP-activated protein kinase gamma 3 subunit (PRKAG3). Detecting
XX the presence of the PRKAG3 DNA, or a variant, is useful in determining
XX a risk estimate of a metabolic disease, such as diabetes or obesity,
XX in a subject. The variation may occur in exons 3, 4 or 10. In exon
XX 3 variation may be a substitution of a G for a C at nucleotide 320,
XX resulting in the amino acid substitution P71A; in exon 4 variation may
XX be a substitution of a T for a C at nucleotide 550; and in exon 10
XX variation may be a substitution of a T for a C at nucleotide 1037,
XX resulting in the amino acid substitution R340W. There may also be
XX nucleotide variation in intron 6. The numbering of these
XX variations is based on the full length cDNA as given, rather than on
XX position 1 of the open reading frame.
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XX Best Local Similarity 100.0%; Pred. No. 1.1e-22;
XX Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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XX 1 ctgtcccgagagcccaattcccaagctggctggatgacgaactcggaaaccggc 60
XX ||||||||||||||||||||||||||||||||||||||||||||||||
XX Db 500 ctgtcccgagagcccaattcccaagctggctggatgacgaactcggaaaccggc 559
XX
XX 61 gcccgatctacatgcgtctcatgcagagcacacctgcta 101
XX ||||||||||||||||||||||||||||||||||||||||||||||||
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XX RESULT 3
XX AAD03296
XX ID AAD03296 standard; DNA; 2109 BP.
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XX AAD03296;
XX
XX 13-JUN-2001 (first entry)
XX
XX Human AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.
XX
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KW human: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
 KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
 KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
 KW cystathione beta synthase; CBS; cardiant; gene therapy; ss.
 OS Homo sapiens.
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 FT 472..1389
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 FT 3'UTR 1390..2109
 FT /*tag= c
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 PD 22-MAR-2001.
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 PD 11-SEP-2000: 2000MO-EP09896.
 PF 10-SEP-1999: 99EP-0402236.
 PR 18-MAY-2000: 2000EP-0401388.
 XX
 PA (INRG) INRA INST NAT RECH AGRONOMIQUE.
 PA (ANDE/) ANDERSSON L.
 PA (LOOF/) LOOFT C.
 PA (KALM/) KALM E.
 XX
 PI Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
 PI Iannucciell N, Gellin J, Le Roy P, Chardon P;
 XX
 DR WPI: 2001-244810/25.
 DR P-PSDB: AAE00221.
 XX
 PT New variants of the gamma subunit of vertebrate adenosine
 PT monophosphate-activated kinase for diagnosis or treatment of disorders
 PT associated with energy metabolism such as diabetes, obesity, and
 PT myopathy -
 XX
 PS Claim 12: Fig 2: 71pp: English.
 XX
 CC The present sequence is a cDNA encoding human adenosine monophosphate
 CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
 CC PRKAG3. Mutation in prkag3 results in an altered regulation of
 CC carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
 CC useful as therapeutic for treating carbohydrate metabolism disorders such
 CC as diabetes, obesity, and disorders associated with muscle metabolism
 CC such as myopathy and cardiovascular diseases, to modulate AMPK
 CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
 CC and its functionally altered mutants are useful for the diagnostic
 CC evaluation, genetic testing and prognosis of a metabolic disorder,
 CC preferably a carbohydrate metabolism disorder. Primers that can detect
 CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
 CC useful for detecting a dysfunction of carbohydrate metabolism resulting
 CC from the expression of a functionally altered allele of PRKAG3.
 CC Transgenic animal and host cell transformed with PRKAG3 or a
 CC heterotimeric AMPK consisting of PRKAG3 or its mutant, are useful for
 CC screening compounds able to modulate AMPK activity. Nucleic acid
 CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
 CC in a sequence encoding the first cystathione beta synthase (CBS) domain
 CC of PRKAG3 and is useful in gene therapy.
 XX
 SQ Sequence 2109 BP: 458 A; 621 C; 560 G; 470 T; 0 other:

Query Match 100.0%; Score 101; DB 22; Length 2109;
 Best Local Similarity 100.0%; Pred. NO. 1,1e-22;
 Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

0Y 1 cgtgcgccgagggccattccccaagctggtcgtggatgcagcaactcgtggaaccgcgc 60
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Db 400 cgtgcgccgagggccattccccaagctggtcgtggatgcagcaactcgtggaaccgcgc 459
 0Y 61 gcccaagatctaacatgcgtcttcacgcagagcaacactgccta 101
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
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RESULT 4
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 ID AAD03320 standard; cDNA: 2115 BP.
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 AC AAD03320;
 XX
 DT 13-JUN-2001 (first entry)
 XX
 DE Human AMPK gamma subunit muscle-specific isoform, complete PRKAG3 cDNA.
 XX
 KW Human: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
 KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
 KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
 KW cystathione beta synthase; CBS; cardiant; gene therapy; ss.
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 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT CDS 1..1395
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 PN MO200120003-A2.
 PD 22-MAR-2001.
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 PD 11-SEP-2000: 2000MO-EP09896.
 PF 10-SEP-1999: 99EP-0402236.
 PR 18-MAY-2000: 2000EP-0401388.
 XX
 PA (INRG) INRA INST NAT RECH AGRONOMIQUE.
 PA (ANDE/) ANDERSSON L.
 PA (LOOF/) LOOFT C.
 PA (KALM/) KALM E.
 XX
 PI Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
 PI Iannucciell N, Gellin J, Le Roy P, Chardon P;
 XX
 DR WPI: 2001-244810/25.
 DR P-PSDB: AAE00223.
 XX
 PT New variants of the gamma subunit of vertebrate adenosine
 PT monophosphate-activated kinase for diagnosis or treatment of disorders
 PT associated with energy metabolism such as diabetes, obesity, and
 PT myopathy -
 XX
 PS Claim 12: Page 65-68: 71pp: English.
 XX
 CC The present sequence is a cDNA encoding human adenosine monophosphate
 CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
 CC complete PRKAG3. Mutation in prkag3 results in an altered regulation of
 CC carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
 CC useful as therapeutic for treating carbohydrate metabolism disorders such
 CC as diabetes, obesity, and disorders associated with muscle metabolism
 CC such as myopathy and cardiovascular diseases, to modulate AMPK
 CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
 CC and its functionally altered mutants are useful for the diagnostic
 CC evaluation, genetic testing and prognosis of a metabolic disorder,
 CC preferably a carbohydrate metabolism disorder. Primers that can detect
 CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
 CC useful for detecting a dysfunction of carbohydrate metabolism resulting
 CC from the expression of a functionally altered allele of PRKAG3.
 CC Transgenic animal and host cell transformed with PRKAG3 or a
 CC heterotimeric AMPK consisting of PRKAG3 or its mutant, are useful for
 CC screening compounds able to modulate AMPK activity. Nucleic acid

CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
 CC in a sequence encoding the first cystathionine beta synthase (CBS) domain
 CC of PRKAG3 and is useful in gene therapy.

SO Sequence 2115 BP: 460 A; 622 C; 562 G; 471 T; 0 other;

Query Match 100.0%; Score 101; DB 22; Length 2115;
 Best Local Similarity 100.0%; Pred. No. 1.1e-22;
 Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ctgtcccgagagcccatctcccaagctggctggatgacgaactgcggaaccgcgc 60
 Db 406 ctgtcccgagagcccatctcccaagctggctggatgacgaactgcggaaccgcgc 60
 OY 61 gccagatctacatgcgcttcacatgcagagacacactgcta 101
 Db 466 gccagatctacatgcgcttcacatgcagagacacactgcta 506

RESULT 5
 AAD03295
 ID AAD03295 standard; cDNA; 1867 BP.
 AC AAD03295;
 XX

13-JUN-2001 (first entry)

Pig AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.

KW genetic testing: obesity; myopathy; cardiovascular disease; anorectic;
 KW genetic testing: carbohydrate metabolism disorder; skeletal muscle;
 KW cystathionine beta synthase; CBS; cardiant; gene therapy; RN locus;
 XX chromosome 15; ss.

Sus scrofa.

Key Location/Qualifiers

FT 5'UTR 1..471

FT CDS /*tag= a 472..1389

FT /*tag= b /product= "Sus scrofa PRKAG3 protein"

FT 3'UTR 1390..1867

FT /*tag= c

WO200120003-A2.

22-MAR-2001.

11-SEP-2000; 2000WO-EP09896.

10-SEP-1999; 99EP-0402236.

18-MAY-2000; 2000EP-0401388.

(INRG) INRA INST NAT RECH AGRONOMIQUE.

(ANDE/) ANDERSSON L.

(LOOF/) LOOFT C.

(KALM/) KALM E.

Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
 Iannucci N, Geil J, Le Roy P, Chardon P;

WT: 2001-244810/25.

P-PSDB: AAE00220.

New variants of the gamma subunit of vertebrate adenosine
 monophosphate-activated kinase for diagnosis or treatment of disorders
 associated with energy metabolism such as diabetes, obesity, and
 myopathy

Claim 12; Fig 2; 71pp; English.

XX The present sequence is a cDNA encoding pig adenosine monophosphate
 CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
 CC PRKAG3. Prkag3 gene is located in the RN locus of chromosome 15.
 CC Mutation in Prkag3 results in an altered regulation of carbohydrate
 CC metabolism, particularly in skeletal muscle. PRKAG3 is useful as
 CC therapeutic for treating carbohydrate metabolism disorders such as
 CC diabetes, obesity, and disorders associated with muscle metabolism
 CC such as myopathy and cardiovascular diseases, to modulate AMPK
 CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
 CC and its functionally altered mutants are useful for the diagnostic
 CC evaluation, genetic testing and prognosis of a metabolic disorder.
 CC Preferably a carbohydrate metabolism disorder. Primers that can detect
 CC a genetic polymorphic marker linked to a sequence encoding PRKAG3,
 CC useful for detecting a dysfunction of carbohydrate metabolism resulting
 CC from the expression of a functionally altered allele of PRKAG3.
 CC Transgenic animal and host cell transformed with PRKAG3 or a
 CC heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
 CC screening compounds able to modulate AMPK activity. Nucleic acid
 CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
 CC in a sequence encoding the first cystathionine beta synthase (CBS) domain
 CC of PRKAG3 and is useful in gene therapy.

SO Sequence 1867 BP: 380 A; 583 C; 529 G; 375 T; 0 other;

Query Match 71.7%; Score 72.4; DB 22; Length 1867;
 Best Local Similarity 83.7%; Pred. No. 1.3e-13;
 Matches 82; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

OY 4 tcccgagagcccatctcccaagctggctggatgacgaactgcggaaccgcgc 63
 Db 403 tcccgagagctgtcgtaccagcagctggctggatgacgaactgcggaaccgcgc 63

OY 64 cagatctacatgcgcttcacatgcagagacacactgcta 101

Db 463 cagatctacatgcgcttcacatgcagagacacactgcta 500

RESULT 6

AAD03319
 ID AAD03319 standard; cDNA; 1873 BP.

AC AAD03319;
 XX

13-JUN-2001 (first entry)

Pig AMPK gamma subunit muscle-specific isoform, complete PRKAG3 cDNA.

KW genetic testing: obesity; myopathy; cardiovascular disease; anorectic;
 KW genetic testing: carbohydrate metabolism disorder; skeletal muscle;
 KW cystathionine beta synthase; CBS; cardiant; gene therapy; RN locus;
 XX chromosome 15; ss.

Sus scrofa.

Key Location/Qualifiers

FT CDS /*tag= a 1..1395

FT /*tag= b /product= "Sus scrofa complete Prkag3 protein"

WO200120003-A2.

22-MAR-2001.

11-SEP-2000; 2000WO-EP09896.

10-SEP-1999; 99EP-0402236.

18-MAY-2000; 2000EP-0401388.

(INRG) INRA INST NAT RECH AGRONOMIQUE.
 (ANDE/) ANDERSSON L.

```

PA (UOFT/) LOOFT C.
PA (KALM/) KALM E.
XX
XX
XX Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
XX Iannuccielli N, Gellin J, Le Roy P, Chardon P;
XX
XX WP1: 2001-244810/25.
XX P-PSDB: AAE00222.
XX
XX New variants of the gamma subunit of vertebrate adenosine
XX monophosphate-activated kinase for diagnosis or treatment of disorders
XX associated with energy metabolism such as diabetes, obesity, and
XX myopathy -
XX
XX Claim 12: Page 62-64; 71pp: English.
XX
XX The present sequence is a cDNA encoding pig adenosine monophosphate
XX (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
XX complete PRKAG3. Prkag3 gene is located in the RN locus of chromosome
XX 15. Mutation in Prkag3 results in an altered regulation of carbohydrate
XX metabolism, particularly in skeletal muscle. PRKAG3 is useful as
XX therapeutic for treating carbohydrate metabolism disorders such as
XX diabetes, obesity, and disorders associated with muscle metabolism
XX such as myopathy and cardiovascular diseases, to modulate AMPK
XX activity, and for restoring a normal AMPK function. PRKAG3 sequence
XX and its functionally altered mutants are useful for the diagnostic
XX evaluation, genetic testing and prognosis of a metabolic disorder,
XX preferably a carbohydrate metabolism disorder. Primers that can detect
XX a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
XX useful for detecting a dysfunction of carbohydrate metabolism resulting
XX from the expression of a functionally altered allele of PRKAG3.
XX Transgenic animal and host cell transformed with PRKAG3 or a
XX heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
XX screening compounds able to modulate AMPK activity. Nucleic acid
XX encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
XX in a sequence encoding the first cystathione beta synthase (CBS) domain
XX of PRKAG3 and is useful in gene therapy.
XX
XX Sequence 1873 BP: 382 A; 580 C; 535 G; 376 T; 0 other;
XX
XX Query Match 71.7%; Score 72.4; DB 22; Length 1873;
XX Best Local Similarity 83.7%; Pred. No. 1,3e-13;
XX Matches 82; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
XX
XX QY 4 tccccgagggcccaattcccaagctgggtggaatgaagaaacccggcgcc 63
XX ||||| ||| | | |||| | ||||| ||||| || |||| ||| || |||
XX Db 409 tccccagaggtgtcttaaccagctgggtggaatgaatgaagaaacccggggcc 468
XX
XX QY 64 cagatctacatgcgtctcatgcagagcaacactgcta 101
XX ||| ||||| |||| | ||||| ||||| ||||| |||||
XX Db 469 caggtctacatgcacttcatgcagagcaacactgcta 506
XX
XX RESULT 7
XX AAD03321
XX ID AAD03321 standard; DNA: 2022 BP.
XX
XX AC AAD03321;
XX
XX XX 13-JUN-2001 (first entry)
XX
XX DE Sus scrofa PRKAG3 splice variant DNA.
XX
XX KW pig; gamma subunit; adenosine monophosphate-activated kinase; AMPK;
XX PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
XX genetic testing; carbohydrate metabolism disorder; skeletal muscle;
XX cystathione beta synthase; CBS; cardiac; gene therapy; ds.
XX
XX OS Sus scrofa.
XX
XX FH Key Location/Qualifiers
XX FT CDS 1..1545

```

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FT /*tag= a
FT /product= "Sus scrofa Prkag3 splice variant"
XX
XX
XX WO200120003-A2.
XX
XX
XX 22-MAR-2001.
XX
XX
XX 11-SEP-2000; 2000MO-EP09896.
XX
XX
XX 10-SEP-1999; 99EP-0402236.
XX 18-MAY-2000; 2000EP-0401388.
XX
XX
XX (INRG ) INRA INST NAT RECH AGRONOME.
XX (ANDE/) ANDERSSON L.
XX (UOFT/) LOOFT C.
XX (KALM/) KALM E.
XX
XX Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
XX Iannuccielli N, Gellin J, Le Roy P, Chardon P;
XX
XX WP1: 2001-244810/25.
XX P-PSDB: AAE00224.
XX
XX New variants of the gamma subunit of vertebrate adenosine
XX monophosphate-activated kinase for diagnosis or treatment of disorders
XX associated with energy metabolism such as diabetes, obesity, and
XX myopathy -
XX
XX Claim 12: Page 69; 71pp: English.
XX
XX The present sequence is pig adenosine monophosphate (AMP)-activated
XX kinase (AMPK) gamma subunit muscle-specific isoform, PRKAG3 splice
XX variant DNA. Prkag3 gene is located in the RN locus of chromosome 15.
XX mutation in Prkag3 results in an altered regulation of carbohydrate
XX metabolism, particularly in skeletal muscle. PRKAG3 is useful as
XX therapeutic for treating carbohydrate metabolism disorders such as
XX diabetes, obesity, and disorders associated with muscle metabolism
XX such as myopathy and cardiovascular diseases, to modulate AMPK
XX activity, and for restoring a normal AMPK function. PRKAG3 sequence
XX and its functionally altered mutants are useful for the diagnostic
XX evaluation, genetic testing and prognosis of a metabolic disorder,
XX preferably a carbohydrate metabolism disorder. Primers that can detect
XX a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
XX useful for detecting a dysfunction of carbohydrate metabolism resulting
XX from the expression of a functionally altered allele of PRKAG3.
XX Transgenic animal and host cell transformed with PRKAG3 or a
XX heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
XX screening compounds able to modulate AMPK activity. Nucleic acid
XX encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
XX in a sequence encoding the first cystathione beta synthase (CBS) domain
XX of PRKAG3 and is useful in gene therapy.
XX
XX Sequence 2022 BP: 412 A; 623 C; 593 G; 394 T; 0 other;
XX
XX Query Match 71.7%; Score 72.4; DB 22; Length 2022;
XX Best Local Similarity 83.7%; Pred. No. 1,3e-13;
XX Matches 82; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
XX
XX QY 4 tccccgagggcccaattcccaagctgggtggaatgaagaaacccggcgcc 63
XX ||||| ||| | | |||| | ||||| ||||| || |||| ||| || |||
XX Db 559 tccccagaggtgtcttaaccagctgggtggaatgaatgaagaaacccggggcc 618
XX
XX QY 64 cagatctacatgcgtctcatgcagagcaacactgcta 101
XX ||| ||||| |||| | ||||| ||||| ||||| |||||
XX Db 619 caggtctacatgcacttcatgcagagcaacactgcta 656
XX
XX RESULT 8
XX AAS51418
XX ID AAS51418 standard; DNA: 1428 BP.
XX
XX AC AAS51418;

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XX 13-FEB-2002 (first entry)
DT
XX
XX Pseudomonas aeruginosa DNA for cellular proliferation protein #3.
DE
XX
XX
XX Antisense: ds: prokaryotic cellular proliferation gene;
KM antibiotic; antibacterial; drug design.
XX
XX
XX Pseudomonas aeruginosa.
OS
XX
XX MO200170955-A2.
PN
XX
XX 27-SEP-2001.
PD
XX
XX
XX 21-MAR-2001: 2001WO-US09180.
PF
XX
XX 21-MAR-2000: 2000US-191078P.
PR 23-MAY-2000: 2000US-206848P.
PR 26-MAY-2000: 2000US-207727P.
PR 23-OCT-2000: 2000US-242578P.
PR 27-NOV-2000: 2000US-253625P.
PR 22-DEC-2000: 2000US-257931P.
PR 16-FEB-2001: 2001US-269308P.
XX
XX (ELITR-) ELITRA PHARM INC.
PA
XX
XX Haselbeck R, Ohlsen KL, Zyskind JM, Wall D, Trawick JD, Carr GJ,
PI Yamamoto RT, Xu HH;
XX
XX WPI: 2001-611495/70.
DR P-PSDB: AAU33559.
XX
XX New polynucleotides for the identification and development of
PT antibiotics, comprise sequences of antisense nucleic acids -
XX
XX
XX Claim 27: Seq ID No 4000: 511pp; English.
PS
XX
XX The invention relates to antisense inhibitors of genes essential to
CC prokaryotic cellular proliferation, their use in identifying the
CC genes, their use in the discovery of novel antibiotics, the essential
CC genes themselves and the encoded proteins. The prokaryotes used are
CC Escherichia coli, Staphylococcus aureus, Salmonella typhi, Klebsiella
CC pneumoniae, Pseudomonas aeruginosa and Enterococcus faecalis. The
CC invention is also useful for the identification of potential new targets
CC for antibiotic development. The antisense nucleic acids can also be used
CC to identify proteins used in proliferation, to express these proteins,
CC and to obtain antibodies capable of binding to the expressed proteins.
CC The proteins can be used to screen compounds in rational drug discovery
CC programmes. The antisense nucleic acid sequence is also useful to screen
CC for homologous nucleic acids which are required for cell proliferation in
CC a wide variety of organisms. The present sequence encodes an
CC essential prokaryotic cellular proliferation protein.
CC Note: The sequence data for this patent did not form part
CC of the printed specification, but was obtained in electronic
CC format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 1428 BP: 206 A: 525 C: 423 G: 274 T: 0 other;
SQ

Query Match 29.9%; Score 30.2; DB 23; Length 1428;
Best Local Similarity 58.2%; Pred. No. 3.2;
Matches 53; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

OY 3 gtcccgagccaccattcccaagctggctggatgacgaatcggaaccggcgc 62
DB 1218 gtccgtatgtgctatatacctgtgacccggcggtgacccgttcattgtgcgcg 1277
OY 63 ccaagatctacatgccttcattcagagagac 93
DB 1278 gctcgagcatcatgctgcatcattcgcggagac 1308

RESULT 9
AAS74198
ID AAS74198 standard; cDNA: 3918 BP.
XX
XX
XX AAS74198:
AC
XX
XX 13-FEB-2002 (first entry)
DT
XX
XX
XX DNA encoding novel human diagnostic protein #10002.
DE
XX
XX Human: chromosome mapping; gene mapping; gene therapy; forensic;
KM food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
XX
XX Homo sapiens.
OS
XX
XX MO200175067-A2.
PN
XX
XX 11-OCT-2001.
PD
XX
XX 30-MAR-2001: 2001WO-US08631.
PF
XX
XX 31-MAR-2000: 2000US-0540217.
PR 23-AUG-2000: 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
PA
XX
XX Drmanac RT, Liu C, Tang YT;
PI
XX
XX WPI: 2001-639362/73.
DR P-PSDB: ABG10011.
XX
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
XX
XX Claim 1: SEQ ID No 10002: 103pp; English.
PS
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 3918 BP: 1328 A: 872 C: 822 G: 896 T: 0 other;
SQ

Query Match 29.1%; Score 29.4; DB 23; Length 3918;
Best Local Similarity 56.8%; Pred. No. 6.9;
Matches 54; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

OY 4 tcccgagccaccattcccaagctggctggatgacgaatcggaaccggcgc 63
DB 2710 tcccttatcccatctccctctcgtcgtggtgacgagcgcggtcgtgcg 2769
OY 64 cagatctacatgccttcattcagagacacactg 98

Db 2770 cagctccacctgcagcccccagctgcagatccactg 2804

RESULT 10

AA580424 ID AAS80424 standard; cDNA; 3918 BP.

AC AAS80424;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #16228.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.

PN WO200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT;

DR WPI: 2001-639362/73.

DR P-PSDB; ABG16237.

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -

Claim 1: SEQ ID No 16228; 103pp; English.

The invention relates to isolated polynucleotide (I) and
polypeptide (II) sequences. (I) is useful as hybridisation probes,
polymerase chain reaction (PCR) primers, oligomers, and for chromosome
and gene mapping, and in recombinant production of (II). The
polynucleotides are also used in diagnostics as expressed sequence tags
for identifying expressed genes. (I) is useful in gene therapy techniques
for restoring normal activity of (II) or to treat disease states involving
quantitating a polypeptide in tissue, as molecular weight markers and as
a food supplement. (II) and its binding partners are useful in medical
imaging of sites expressing (II). (I) and (II) are useful for treating
disorders involving aberrant protein expression or biological activity.
The polypeptide and polynucleotide sequences have applications in
diagnostics, forensics, gene mapping, identification of mutations
and to produce other types of data and products dependent on DNA and
amino acid sequences. AAS64197-AAS94564 represent novel human
diagnostic coding sequences of the invention.
Note: The sequence data for this patent did not appear in the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences.

Sequence 3918 BP: 1328 A; 872 C; 822 G; 896 T; 0 other;

Query Match 29.1%; Score 29.4; DB 23; Length 3918;
Best Local Similarity 56.8%; Pred. No. 6.9;

Matches 54; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

OY 4 tcccccagagcccatcctccaaagctggcgtggaagaagcagcgaacccgcgcgc 63
DB 2710 tccctatcccccacatccctcctgctgaggtgtgtgtggcgacagcgcggaactgcgcgg 2769

OY 64 cagatctacatgcgtctcatgcagagacacactg 98
DB 2770 cagctccacctgcagcccccagctgcagatccactg 2804

RESULT 11

AA574446 ID AAS74446 standard; cDNA; 4397 BP.

AC AAS74446;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #10250.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.

PN WO200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT;

DR WPI: 2001-639362/73.

DR P-PSDB; ABG10259.

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -

Claim 1: SEQ ID No 10250; 103pp; English.

The invention relates to isolated polynucleotide (I) and
polypeptide (II) sequences. (I) is useful as hybridisation probes,
polymerase chain reaction (PCR) primers, oligomers, and for chromosome
and gene mapping, and in recombinant production of (II). The
polynucleotides are also used in diagnostics as expressed sequence tags
for identifying expressed genes. (I) is useful in gene therapy techniques
for restoring normal activity of (II) or to treat disease states involving
quantitating a polypeptide in tissue, as molecular weight markers and as
a food supplement. (II) and its binding partners are useful in medical
imaging of sites expressing (II). (I) and (II) are useful for treating
disorders involving aberrant protein expression or biological activity.
The polypeptide and polynucleotide sequences have applications in
diagnostics, forensics, gene mapping, identification of mutations
and to produce other types of data and products dependent on DNA and
amino acid sequences. AAS64197-AAS94564 represent novel human
diagnostic coding sequences of the invention.
Note: The sequence data for this patent did not appear in the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences.

Sequence 4397 BP: 1506 A; 980 C; 935 G; 976 T; 0 other;

Query Match 29.1%; Score 29.4; DB 23; Length 4397;
Best Local Similarity 56.8%; Pred. No. 7;

Matches 54; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

OY 4 tcccgagagcccatlcccaagctgggtgagatgacgaactgcgaaaccggcgc 63
 Db 3189 tccctatcccatlcccatlccctccctgaggtgtgtggtgcacagcgaggtgcg 3248
 OY 64 cagatctacatgcgtctcatgacgagacacactg 98
 Db 3249 cagctccactgcagcccgagtgagatccactg 3283

RESULT 12

AAS73340 ID AAS73340 standard: cDNA; 5122 BP.

AC AAS73340:

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #9144.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.

PN WO200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT;

DR WPI: 2001-639362/73.

DR P-PSDB: ABC09153.

PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -

PS Claim 1; SEQ ID NO 9144; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 5122 BP; 1815 A; 1151 C; 1032 G; 1124 T; 0 other;

Query Match

29.1%; Score 29.4; DB 23; Length 5122;

Best Local Similarity 56.8%; Pred. No. 7.2;
 Matches 54; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

OY 4 tcccgagagcccatlcccaagctgggtgagatgacgaactgcgaaaccggcgc 63
 Db 3914 tccctatcccatlcccatlccctccctgaggtgtgtggtgcacagcgaggtgcg 3973

OY 64 cagatctacatgcgtctcatgacgagacacactg 98
 Db 3974 cagctccactgcagcccgagtgagatccactg 4008

RESULT 13

ABL19505 ID ABL19505 standard: DNA; 1406 BP.

AC ABL19505;

DT 26-MAR-2002 (first entry)

DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 9988.

KW Drosophila; developmental biology; cell signalling; insecticide;
 KW pharmaceutical; gene; ds.

OS Drosophila melanogaster.

PN WO200171042-A2.

PD 27-SEP-2001.

PF 23-MAR-2001; 2001WO-US09231.

PR 23-MAR-2000; 2000US-191637P.

PR 11-JUL-2000; 2000US-0614150.

PA (PEKE) PE CORP NY.

PI Venter JC, Adams M, Li PWD, Myers EW;

DR WPI: 2001-656860/75.

PT New isolated nucleic acid detection reagent for detecting 1000 or more
 PT genes from Drosophila and for elucidating cell signalling and cell-cell
 PT interactions -

PS Claim 1; SEQ ID NO 9988; 21pp + Sequence Listing; English.

XX The invention relates to an isolated nucleic acid detection reagent
 CC capable of detecting 1000 or more genes from Drosophila. The invention is
 CC useful in developmental biology and in elucidating cell signalling and
 CC cell-cell interactions in higher eukaryotes for the development of
 CC insecticides, therapeutics and pharmaceutical drugs. The invention
 CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
 CC sequences (ABL01840-ABL16175) and the encoded proteins
 CC (ABB57737-ABB72072).
 CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 1406 BP; 358 A; 409 C; 404 G; 235 T; 0 other;

Query Match

Best Local Similarity 71.7%; Score 29; DB 23; Length 1406;
 Matches 38; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

OY 49 cggaaaccggcgccagatctacatgcgtctcatgacgagacacactgcta 101
 Db 264 ctgaagccggcgccgctgtgcccgcgctgttcaaggagctctccgacta 316

RESULT 14

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ABL19504
ID ABL19504 standard; DNA: 3406 BP.
XX
AC ABL19504;
XX
XX
XX 26-MAR-2002 (first entry)
XX
DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 9985.
XX
XX Drosophila: developmental biology; cell signalling; insecticide;
XX
XX pharmaceutical; gene; ds.
XX
OS Drosophila melanogaster.
XX
XX WO200171042-A2.
XX
XX 27-SEP-2001.
XX
XX 23-MAR-2001: 2001WO-US09231.
XX
XX 23-MAR-2000: 2000US-191637P.
XX
XX 11-JUL-2000: 2000US-0614150.
XX
XX (PEKE ) PE CORP NY.
XX
XX Venter JC, Adams M, Li PWD, Myers EW;
XX
XX WPI: 2001-656860/75.
XX
XX New isolated nucleic acid detection reagent for detecting 1000 or more
XX genes from Drosophila and for elucidating cell signalling and cell-cell
XX interactions -
XX
XX Claim 1: SEQ ID NO 9985; 21pp + Sequence Listing: English.
XX
XX The invention relates to an isolated nucleic acid detection reagent
XX capable of detecting 1000 or more genes from Drosophila. The invention is
XX useful in developmental biology and in elucidating cell signalling and
XX cell-cell interactions in higher eukaryotes for the development of
XX insecticides, therapeutics and pharmaceutical drugs. The invention
XX discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
XX sequences (ABB57737-ABB72072).
XX
XX The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 3406 BP; 902 A; 852 C; 828 G; 824 T; 0 other;
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Query Match 28.7%; Score 29; DB 23; Length 3406;
Best Local Similarity 71.7%; Pred. No. 9;
Matches 38; Conservative 0; Mismatches 15; Indels 0; Gaps 0;
OY 49 cggaaacccggcgccagatctacatgcttcacgagagacacactgcta 101
DB 1264 ctgaagccggcgccgctgtccgcgctgttcacgagagctctccgacta 1316
RESULT 15
ABL19502
ID ABL19502 standard; DNA: 3593 BP.
XX
XX
XX ABL19502;
XX
XX 26-MAR-2002 (first entry)
XX
XX Drosophila melanogaster genomic polynucleotide SEQ ID NO 9979.
XX
XX Drosophila: developmental biology; cell signalling; insecticide;
XX pharmaceutical; gene; ds.
XX
OS Drosophila melanogaster.

```

```

XX
XX PN WO200171042-A2.
XX
XX PD 27-SEP-2001.
XX
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XX 23-MAR-2001: 2001WO-US09231.
XX
XX 23-MAR-2000: 2000US-191637P.
XX
XX 11-JUL-2000: 2000US-0614150.
XX
XX (PEKE ) PE CORP NY.
XX
XX Venter JC, Adams M, Li PWD, Myers EW;
XX
XX WPI: 2001-656860/75.
XX
XX New isolated nucleic acid detection reagent for detecting 1000 or more
XX genes from Drosophila and for elucidating cell signalling and cell-cell
XX interactions -
XX
XX Claim 1: SEQ ID NO 9979; 21pp + Sequence Listing: English.
XX
XX The invention relates to an isolated nucleic acid detection reagent
XX capable of detecting 1000 or more genes from Drosophila. The invention is
XX useful in developmental biology and in elucidating cell signalling and
XX cell-cell interactions in higher eukaryotes for the development of
XX insecticides, therapeutics and pharmaceutical drugs. The invention
XX discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
XX sequences (ABB57737-ABB72072).
XX
XX The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 3593 BP; 920 A; 953 C; 902 G; 818 T; 0 other;
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Query Match 28.7%; Score 29; DB 23; Length 3593;
Best Local Similarity 71.7%; Pred. No. 9;
Matches 38; Conservative 0; Mismatches 15; Indels 0; Gaps 0;
OY 49 cggaaacccggcgccagatctacatgcttcacgagagacacactgcta 101
DB 3377 ctgaagccggcgccgctgtccgcgctgttcacgagagctctccgacta 3429

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Search completed: October 3, 2002, 16:30:46
 Job time: 14316 sec



GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:15:04 ; Search time 5701.1 Seconds
(without alignments)
239.110 Million cell updates/sec

Title: US-09-826-581-5_COPY_500_600
Sequence: 101
1 cgtgcccgagagccatc.....atgcagagagacacccgcta 101

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : EST:*

1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estnu:*
5: em_estow:*
6: em_estpl:*
7: em_estro:*
8: em_hlc:*
9: gb_est1:*
10: gb_est2:*
11: gb_hlc:*
12: gb_gss:*
13: em_gss_hum:*
14: em_gss_inv:*
15: em_gss_pln:*
16: em_gss_vrt:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	72.2	71.5	444	10	BF890374 291826 MA
2	72.2	71.5	548	10	BI775360 467815 MA
3	67.6	66.9	685	9	BB630381 BB630381
4	31.4	31.1	326	9	AI106144
5	30.6	30.3	679	10	BE340087
6	30.4	30.1	699	10	BE340087 601060879
7	30.2	29.9	580	12	BI199916 602760950
8	30.2	29.9	894	12	BH140228 ZMMBB000
9	29.8	29.5	910	12	CNS01MA1
10	29.8	29.5	1010	12	CNS02SYT
11	29.6	29.3	655	10	CNS03856
12	29.6	29.3	930	10	BM133818
13	29.4	29.1	955	10	BE269692
14	29.2	28.9	1000	10	BE269692 601185836
15	29.2	28.7	313	9	BM468546
16	29.2	28.7	436	10	AA264931
17	29.2	28.7	546	9	AA264931

18	29	28.7	554	9	A1260567
19	29	28.7	561	10	B1229683
20	29	28.7	583	10	B1232284
21	29	28.7	589	9	AA539970
22	29	28.7	627	10	B1633071
23	29	28.7	629	9	A1259247
24	29	28.7	666	10	B1242279
25	29	28.7	738	9	A1544103
26	28.8	28.5	569	10	BG414376
27	28.8	28.5	750	10	BM019844
28	28.6	28.3	339	10	BF773519
29	28.6	28.3	445	10	BC558441
30	28.6	28.3	448	9	AM497449
31	28.6	28.3	509	10	BC560394
32	28.6	28.3	591	9	AM678584
33	28.6	28.3	593	12	BH140663
34	28.6	28.3	605	9	AA195150
35	28.6	28.3	707	12	AG130442
36	28.6	28.3	718	9	AV753229
37	28.6	28.3	742	10	BE394076
38	28.6	28.3	904	10	BF033403
39	28.4	28.1	260	9	AM816448
40	28.4	28.1	712	9	BB623242
41	28.4	28.1	812	9	AA519383
42	28.2	27.9	517	9	AM631272
43	28.2	27.9	525	9	AA922163
44	28.2	27.9	592	12	AZ870322
45	28.2	27.9	802	12	BH077939

ALIGNMENTS

RESULT 1
BF890374 444 bp mRNA linear EST 25-Apr-2001
LOCUS 291826 MARC 3BOV Bos taurus cDNA 5', mRNA sequence.
DEFINITION BF890374
ACCESSION BF890374
VERSION BF890374.1 GI:12281760
KEYWORDS EST.
SOURCE COW.
ORGANISM Bos taurus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovinae; Bos.

REFERENCE

1 (bases 1 to 444)
Smith,T.P.L., Grose,W.M., Freking,B.A., Roberts,A.J., Stone,R.T.,
Casas,E., Wray,J.E., White,J., Cho,J., Fahrenkrug,S.C., Bennett,
G.L., Heaton,M.P., Laegreid,W.M., Rohrer,G.A., Chitko-McKown,C.G.,
Peters,G., Holt,I., Karaycheva,S., Liang,F., Quackenbush,J. and
Keefe,J.W.

Sequence evaluation of four pooled-tissue normalized bovine CDNA
libraries and construction of a gene index for cattle
Genome Res. 11 (4), 626-630 (2001)
21180013

COMMENT

USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt_c trimmed with phred
v0.980904.e. Vector identified by cross-match with the -minscore 18
and -mismatch 12 options.

PCR Primers

FORWARD: AGCAACACCTATGACAT
BACKWARD: GTTCCACGTCACGACG
Plate: 57 row: H column: 11
Seq primer: ATTACGCTACCTATAG.

FEATURES

1..444
/organism="Bos taurus"
/db_xref="taxon:9913"

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/cclone_1lb="MARC 3BOV"
/tissue_type="pooled"
/lab_host="DH10B"
/notes="Vector: pCMV Sport6; Site_1: XbaI; Site_2: XhoI;
library made from pooled tissue from marrow, alveolar
macrophage, ovary, fetal semitendinosus muscle, and fetal
longissimus muscle."
BASE COUNT      91 a      147 c      139 g      67 t
ORIGIN

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Query Match	71.5%	Score 72.2	DB 10	Length 444
Best Local Similarity	82.2%	Pred. No. 7.5e-11		
Matches 83	Conservative	0	Mismatches 18	Indels 0
			Gaps	0

Oy	1	cctgcaccccgagggcccatttccccaaactcgtgcatgtacgaacatcgagaaaccggc	60
Db	295	CCGTCGCCCGCAGGGTGGCGGTGCCTCACGGCTGTGGCGTGACATGACGTGGGAACCCAGGG	350
Oy	61	gccacagatacatactgcttcataatgcaagaagacaaccttgcta	101
Db	355	GCTCAGCGCTACATGACAACTTTCAATCATCAGAGAACACACTGGTCTA	395

RESULT 2	LOCUS	548 bp	mRNA	linear	EST 25-SEP-2001
B1775360	B1775360				
DEFINITION	467815 MARC 2BOV Bos taurus cDNA 5', mRNA sequence.				

KEYWORDS
SOURCE

ORGANISM	REFERENCE
Bos taurus	Smith, T. P. L., Grosse, W. M., Freking, B. A., Roberts, A. J., Stone, R. T.
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;	
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;	
Bovidae; Bovinae; Bos.	
1 (bases 1 to 548)	

TITLE
Sequence evaluation of four pooled-tissue normalized bovine cDNA
libraries and construction of a gene index for cattle
JOURNAL
Genome Res. 11 (4), 626-630 (2001)

JOURNAL	Genome Res. 11 (4), 626-630 (2001)
MEDLINE	21180013
COMMENT	Contact: Smith TPL

Email: sm1the@mail.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred v0.980904.e. Vector identified by cross_match with the -mismatch 12 options.
PCR primers
FORWARD: AGGAAACAGCTATGACCAT
BACKWARD: GTTTCACAGTCACGACG
Plate: 90 row: 1 column: 16
Seq primer: ATTAGGTGACACTATAG.

FEATURES
SOURCE

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/organism="Bos taurus"
/db_xref="taxon:9913"
/clone_lib="MARC 2BOV"
/tissue_type="pooled"

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BASE COUNT	119 a	173 c	163 g	93 t
ORIGIN				

Query Match	71.5%	DB 10	Length 548
Best Local Similarity	82.2%	Pred. No. 7.9e+11	
Matches 83, Conservative	0	Mismatches 18	Indels 0: Gaps 0

Dy 61 gccacagatctcacatgcgtccttcatacgagagacacacctgta 10b
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 397 gctcagctcacatgcacttcatgcagagacacacctgcta 437

LOCUS	DEFINITION	EST
BB630381	665 bp, mRNA	26-OCT-2001
BB630381	RIKEN full-length enriched, 6 days neonate skin Mus	
musculus	CDNA clone A030014A04 5', mRNA sequence.	

SOURCE ORGANISM	house mouse Mus musculus
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3	3
4	4
5	5
6	6
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100	100

REFERENCE
AUTHORS
Atakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanaigaki, T., Hara, A.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 685)

AUTHORS
Arikawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T., Hata, A.,
Himeno, K., Horii, F., Ishii, Y., Ito, M., Kawaj, J., Kono, H., Kouda, K.,
M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K., Ohno, M.,
Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasak, K.,
D., Shibata, K., Shingawa, A., Shiraki, T., Sogabe, Y., Suzuki, H.,
Tagami, M., Tagawa, A., Takahashi, F., Takeda, Y., Tanaka, T., Toyata, T.,
Muramatsu, M. and Hayashizaki, Y.
RIKEN MOUSE ESTS (Arikawa, T., et al. 2001)

TITLE	RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
JOURNAL	Unpublished (2001)
COMMENT	Contact: Yoshinori Hayashizaki

1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-res@gsr.riken.go.jp,
URL: <http://genome.qsc.riken.go.jp/>

Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Kono, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. *Genome Res.* 10 (10), 1617-1630 (2000)

Wagil, K., Fujiwake, S., Inoue, K., Togawa, Y., Itawa, M., Ohara, E., Matshiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsunaga, S., Kawai, U., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.

Kono, H., Fukunishi, Y., Shiba, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y. (2000) Riken integrated sequence analysis (RISA) system—334-Format sequencing pipeline with 384 multicapillary sequencer. *Genome Res* 10 (11), 1757–1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)

Kondo, S., Shingawa, A., Saito, T., Kiyosawa, H., Yamane, I., Mizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and Hayashizaki, Y.

Computational Analysis of full-length mouse cDNAs compared with Human Genome Sequences. *Mamm. Genome*. 12, 673-677 (2001). Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.
e mouse tissues.

FEATURES	Location/Qualifiers
source	1. .685

FEATURES	source
Query Match	66.9% Score 67.6; DB 9; Length 685;
Best Local Similarity	80.6%; Prod. No. 1.7e-09;
Matches 79; Conservative	0; Mismatches 19; Indels 0; Gaps 0;
Db	4
LOCUS	A1106144/C
DEFINITION	db03dl2_p1 2F adult heart library Danio rerio cDNA 5 prime similar
ACCESSION	A1106144
VERSION	A1106144.1
KEYWORDS	EST.
SOURCE	zebrafish.
ORGANISM	Danio rerio.
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio.
AUTHORS	Chen, J.N., Desauvage, F., Hosobuchi, M., Jackson, D.G. and Fishman, M.C.
TITLE	Expressed Sequences from The Adult Zebrafish Heart
JOURNAL	Unpublished (1998)
COMMENT	Contact: Mark C. Fishman Cardiovascular Research Center Massachusetts General Hospital Mail code 1494100A, 149 13th Street, Charlestown, MA 02129, USA Fax: 617265806 Email: fishman@cvc.harvard.edu http://zebrafish.mgh.harvard.edu The original clones used for sequencing are no longer available; the library is available from Mark C. Fishman. Insert length: 326 Std Error: 0.00 Seq primer: p1. Location/Qualifiers 1..326

BASE COUNT	83 a	66 c	96 g	79 t	2 others
ORIGIN					
Query Match	31.1%: Score 31.4; DB 9; Length 326;				
Best Local Similarity	62.7%: Pred. No. 34;				
Matches	47; Conservative 0; Mismatches 28; Indels 0; Gaps 0;				
Qy	23	ccaagctggagctggagatgcagcaactgcggaacccgcgcgccagatctactatgcgtctca	82		
Db	256	CCAAACTGGAAGGCTCTTGATGTCCTNCACAGACTCGCGGACGATCAACAGATTGCTCTCA	197		
Qy	83	tgcaaggagcacacct	97		
Db	196	GGCAGGCTCACACCT	182		
RESULT	5	BE540087	679 bp	MRNA	linear
LOCUS	601060879P2 NIH_MGC_10 Homo sapiens cDNA clone IMAGE:3447378 5',				
DEFINITION	MRNA sequence.				
ACCESSION	BE540087				
VERSION	BE540087.1 GI:9768732				
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniota; Vertebrata; Euteleostomi;				
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.				
TITLE	NIH-MGC http://mgc.nci.nih.gov/.				
JOURNAL	National Institutes of Health, Mammalian Gene Collection (MGC)				
COMMENT	Unpublished (1999)				
	Contact: Robert Strausberg, Ph.D.				
	Email: cgapbs-remail.nih.gov				
	Tissue Procurement: ATCC				
	cDNA Library Preparation: Life Technologies, Inc.				
	cDNA Library Arrayed by: Incyte Genomics, Inc.				
	DNA Sequencing by: Incyte Genomics, Inc.				
	Clone distribution: MGC clone distribution information can be				
	found through the I.M.A.G.E. Consortium/LINL at:				
	http://image.llnl.gov				
	Plate: LAMB420 row: n column: 19				
	High quality sequence stop: 668.				
	Location/Qualifiers				
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	/clone_lib="NIH_MGC_10"				
	/cell_line="MGC36"				
	/lab_host="DH10B"				
	/note="Organ: cervix; Vector: pCMV-SPORT6; Site:1; NotI;				
	Site:2; SalI; Cloned unidirectionally. Primer: Oligo dT.				
	Average insert size 1.5 kb. Library prepared by Life				
	Technologies."				
BASE COUNT	124 a	226 c	177 g	152 t	
ORIGIN					
Query Match	30.3%: Score 30.6; DB 10; Length 679;				
Best Local Similarity	62.3%: Pred. No. 68;				
Matches	48; Conservative 0; Mismatches 29; Indels 0; Gaps 0;				

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Oy 9 gcagggcccatctcccaagctggagatgacgaactcggaaccggcgccagat 68
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Db 244 GCACCCCGCTTTCTGTGAGGTCTGCAGACAGCAGACTGGGACAGCTGGCTGCAT 185
Oy 69 ctacatgcgtctcatgc 85
    || | | | | | |
Db 184 TCACCTGCACATACATGC 168

RESULT 6
B1199916/c 699 bp mRNA linear EST 10-JUL-2001
LOCUS B1199916 602760950F1 NIH_MGC_19 Homo sapiens cDNA clone IMAGE:4896324 5',
DEFINITION mRNA sequence.
ACCESSION B1199916
VERSION B1199916.1 GI:14654937
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota: Chordata: Craniata: Vertebrata: Euteleostomi:
Mammalia: Eutheria: Primates: Catarrhini: Homnidae: Homo.
REFERENCE NIH-MGC http://mgc.ncl.nih.gov/
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabs-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at: image.lnl.gov
Plate: LNCMT783 row: c column: 13
High quality sequence start: 5
High quality sequence stop: 669.
Location/Qualifiers
SOURCE
1. .699
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/db_xref="taxon:9606"
/clone="IMAGE:4896324"
/clone_1b="NIH_MGC_19"
/tissue_type="neuroblastoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: Brain; Vector: pORF7; site:1: XhoI; site:2:
EcoRI; cDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(C). Library constructed by Ling Hong
in the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-CDNA synthesis kit
(stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library."
BASE COUNT 120 a 216 c 222 g 141 t
ORIGIN

```

```

DEFINITION ZM8BB0001118f Maize B73 Zea mays genomic clone ZM8BB0001118f, DNA
sequence.
ACCESSION B1140228
VERSION B1140228.1 GI:1509289
KEYWORDS GSS.
SOURCE Zea mays.
ORGANISM Zea mays
Eukaryota: Viridiplantae: Streptophyta: Embryophyta: Tracheophyta:
Spermatophyta: Magnoliophyta: Liliopsida: Poales: Poaceae: PACC
clade: Panicoideae: Andropogoneae: Zea.
REFERENCE Tomkins,J.P., Main,D., Golceachea,J.L., Frisch,D.A. and Wing,R.A.
AUTHORS A Deep-Coverage BAC Library for Maize
JOURNAL Unpublished (2001)
COMMENT Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: twing@clemson.edu
Seq primer: TAATGACTCCTATAGCG
Class: BAC ends
High quality sequence start: 71
High quality sequence stop: 510.
Location/Qualifiers
SOURCE
1. .580
/organism="Zea mays"
/strain="B73"
/cultivar="B73"
/db_xref="taxon:4577"
/clone="ZM8BB0001118f"
/clone_1b="Maize B73"
/tissue_type="Young leaves"
/lab_host="E. coli"
/note="Vector: pCUGIBAC-1; site:1: HindIII; site:2: NotI;
For more details on library preparation, ordering clones
and sequence analysis see
http://www.genome.clemson.edu/projects/stc/maize/ZM8BB"
BASE COUNT 74 a 221 c 138 g 146 t
ORIGIN

```

```

Query Match 29.9%: Score 30.2: DB 12: Length 580:
Best Local Similarity 62.7%: Pred. No. 85:
Matches 47: Conservative 0: Mismatches 28: Indels 0: Gaps 0:
Oy 27 gctgggctggatgacgaactcggaaccggcgccagatctacatgcgtctatgca 86
    || | ||||| || | ||||| || | ||||| || | || ||||| || |
Db 147 GCCGACTCGGACGACAACTGAGAGAGCGGCTCGAGGGGCTCCACCGCAGTAGCA 88
Oy 87 ggaagcacacctgcta 101
    | | | | | | |
Db 87 GTTTCGACCGGCGA 73

RESULT 8
CNS01M1/c 894 bp DNA linear GSS 14-JUN-2001
LOCUS CNS01M1 Anopheles gambiae GSS SP6 end of clone 21E19 of NotreDame1 library
DEFINITION from strain PEST of Anopheles gambiae (African malaria mosquito),
genomic survey sequence.
ACCESSION AL150666
VERSION AL150666.1 GI:7011145
KEYWORDS GSS.
SOURCE African malaria mosquito.
ORGANISM Anopheles gambiae
Eukaryota: Metazoa: Arthropoda: Tracheata: Hexapoda: Insecta:
Pterygota: Neoptera: Endopterygota: Diptera: Nematocera:
Culicoidae: Anopheles.
REFERENCE 1 (bases 1 to 894)
AUTHORS Genoscope.
TITLE Direct Submission

```

REFERENCE	TITLE	JOURNAL	AUTHORS	COMMENT
Submitted (16-FEB-2000)	Genoscope - Centre National de Sequencage			
BP 191 91006 EVRY cedex - FRANCE (E-mail : sequef@genoscope.cns.fr				
- Web : www.genoscope.cns.fr)				
2 (bases 1 to 894)				
Roeth,C.W., Brey,P.T., Ke,Z., Collins,F.H. and Weissenbach,J.				
Direct Submission				
Submitted (16-FEB-2000)	BBM1, Institut Pasteur, 25, rue du Dr.			
Roux, Paris 75015, France				
This clone is from an A. gambiæ BAC library provided by F.H.				
Collins and sequenced by Genoscope in collaboration with the				
Laboratory of Biochem. and Biol. Molec. of Insects, Institut				
Pasteur.				
location/Qualifiers				
1. 894				
/organism="Anopheles gambiae"				
/strain="PEST"				
/db_xref="taxon:7165"				
/clone="21E19"				
/clone.lib="Notredame1"				
/note="end : SP6"				
HASH COUNT	177 a	256 c	234 g	225 t
OTHERS				2 others
Query Match	29.9%	Score 30.2:	DB 12:	Length 894:
Best Local Similarity	62.7%	Pred. No. 94:		
Matches	47:	Conservative	0:	Mismatches 28: Indels 0: Gaps 0:
Oy	27	gctggcgtggatgacgaactcgcgaaacccgcgcacagatcatcagcttcatacga	86	
Db	181	gctgtgggtgattacgtctctcgtccgccgcacacgattccacacgggtttcttcca	122	
Oy	87	ggagcagaccctgcta	101	
Db	121	ggagcagaccctgcta	107	
RESULT	9			
CNS02SYT		910 bp	DNA	linear
LOCUS				
DEFINITION				
Accession				
VERSION				
KEYWORDS				
SOURCE				
ORGANISM				
REFERENCE				
TITLE				
JOURNAL				
AUTHORS				
COMMENT				

```

FEATURES      http://www.genoscope.cns.fr/Tetraodon.
               Location/Qualifiers
               1..910
               /organism="Tetraodon nigroviridis"
               /db_xref="taxon:998083"
               /clone="163617"
               /clone_11b="G"
               /note="Genoscope sequence ID : COAG163AD09LPI-end : 17"
BASE COUNT    179 a      263 c      305 g      160 t
ORIGIN

```

[illegible]

RESULT	10		
CNS03856			
LOCUS			
DEFINITION	CNS03856	1010 bp	DNA
	Tetraodon nigroviridis genome survey sequence		linear
	005M22 of library G from Tetraodon nigroviridis, genomic survey		17 end of clone
	sequence.		
ACCESSION	AL233007		
VERSION	AL233007.1	GI:7892142	
KEYWORDS	GSF: genome survey sequence.		
SOURCE	Tetraodon nigroviridis		
ORGANISM	Tetraodon nigroviridis		

REFERENCE	AUTHORS	TITLE	JOURNAL	REFERENCE	AUTHORS	TITLE	JOURNAL
1	Roest-Crollius,H., Jalllon,O., Dasilva,C., Fitzames,C., Fisher,C., Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and Weissenbach,J.	Characterization and repeat analysis of the freshwater puterfish Tetraodon nigroviridis	unpublished	2	Roest-Crollius,H., Jalllon,O., Dasilva,C., Bouneau,L., Fisher,C., Bernot,A., Fitzames,C., Wincker,P., Brottler,P., Quetier,F., Saurin,W. and Weissenbach,J.	Human gene number estimate provided by genome wide analysis using Tetraodon nigroviridis DNA sequence	unpublished
3	(bases 1 to 1010)			3	(bases 1 to 1010)		
	Genoscope.				Genoscope.		
	Direct Submission				Submitted (12-Apr-2000) to the EMBL/GenBank/DBJ databases		
	This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetraodon nigroviridis genome. For more information, please take a look at http://www.genoscope.cns.fr/tetraodon .				location/Qualifiers		
	1..1010						

BASE COUNT	/note="Genoscope sequence ID : COBG005DGI11XD1-end : T7"			
ORIGIN	174 a	313 c	312 g	210 t
				1 others

	Query Match	Best Local Similarity	Matches	Conservative	Score	DB 12	length	Gaps
Oy	30	99.9%	43	0	29.8	DB 12	1010	0
	gggctggatagacgaactcggaacccgcgcgcacagatctacatgcttatacaga						89	
Db	620	ggcgatgaaagcgttccacacgtacccccaagcacttcgacttccggacatcagaca					679	
Oy	90	99.9%	43	0	29.8	DB 12	1010	0
	ggcaca						94	
Db	680	gcaca					684	

RESULT	11
LOCUS	BM133818
DEFINITION	BM133818 655 bp mRNA linear EST 07-JAN-2002
LOCUS	NX1LV_012_F09_F NX1LV (Nsif Xylem Late wood vertical) Pinus taeda CDNA clone NX1LV_012_F09 5', mRNA sequence.
ACCESSION	BM133818
VERSION	BM133818.1
KEYWORDS	GI:17141057
SOURCE	EST.
ORGANISM	loblolly pine.
	Pinus taeda
	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Coniferopsida; Coniferales; Pinaceae; Pinus; Pinus.
REFERENCE	1 (bases 1 to 655)
AUTHORS	Sedgeroff, R.
TITLE	Molecular Basis of Wood Formation in the Pine Megagenome
JOURNAL	Unpublished (2000)
COMMENT	Contact: Johnson, Arthur

Seq primer: T3.
Location/Qualifiers
1. .655
/organism="Pinus taeda"
/strain="Coastal plain loblolly pine from North Carolina"
/db_xref="taxon:3352"
/clone="NXLV.012.F09"
/clone_id="NXLV (NsI xylem Late wood Vertical)"
/tissue_type="primary xylem"
/dev_stage="late wood"
/lab_host="XL1-Blue"
/note="Vector: pRipLEX; Site_1: EcoRI: The library is from late (summer-August) wood, taken from below the crown of a 20 year old tree. The harvested xylem tissue was on the cusp between transitional and mature wood. NOTE: The sequences contain a 'cDNA adapter' between the EcoRI site and the start of the EST. The adapter sequence is 'AATTCGCCCATTAATGCC'."

	Query Match	29.3%;	Score 29.6;	DB 10;	Length 655;
	Best Local Similarity	57.6%;	Pred. No. 1.3e+02;		
	Matches 53; Conservative	0;	Mismatches 39;	Indels 0;	Gaps 0;
Oy	4	tcccccagaggccccatttcccaagaacttggtgcggaatagacaactcgtcggaaccggcgac	63		
Db	235	ttcccgaggaaaccaattttccagAACCAAGGTATGATGCCCAACCGCCGCACCCCTAATCA	294		
Oy	64	cagatctacatagtgccttcacatgatgagagacac	95		
Db	295	CAGTTTCATCATGCCCATCTACAATTTCACAC	326		
RESULT 12					
LOCUS	BE269692		930 bp	mRNA	linear EST 13-JUL-2000
DEFINITION	6011185836f1 NIH_MCC_8 Homo sapiens cDNA clone IMAGE:3543584 5'				

ACCESSION	BE269692	BE269692.1	GI:9143317
VERSION	EST.		
KEYWORDS			
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Ekaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi: Mammalia: Eutheria: Primates: Catarrhini: Hominiidae: Homo. 1 (bases 1 to 930)		
AUTHORS	NIH-MGC	http://mgc.ncl.nih.gov/	
TITLE	National Institutes of Health, Mammalian Gene Collection (MGC)		
JOURNAL	Unpublished (1999)		
COMMENT	Contact: Robert Strausberg, Ph.D. Email: ccgabs-remail.nih.gov		
	Tissue Procurement: Louis M. Staudt, M.D., Ph.D.		
	CDNA Library Preparation: Ling Hong/Rubin Laboratory		
	CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)		
	Clone Sequencing by: Incyte Genomics, Inc.		
	Gene distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: image.llnl.gov		
	Plate: LHCW37 row: 9 column: 09		
	High quality sequence stop: 238.		
	Localization: cytoplasmic		

```

FEATURES
source
Location/Qualifiers
1. .930
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3543584"
/clone_id="NH_MGC_8"
/tissue_type="Burkitt lymphoma"
/lab_host="DH10B (phage-resistant)"
/name="Organ: lymph; Vector: pOTB7; site:1: XhoI; site_2: EcoRI; CDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using Zap-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies).".
BASE COUNT
226 a 228 c 267 g 209 t
ORIGIN

```

RESULT	13
LOCUS	BG394974
DEFINITION	602457389F1 NIH_MGC_16 Homo sapiens cDNA clone IMAGE:579696 5', mRNA sequence.
ACCESSION	BG394974
VERSION	BG394974.1 GI:13288422
KEYWORDS	EST.
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
TITLE	1 (bases 1 to 955)
JOURNAL	NIH-MGC http://mgc.nci.nih.gov/ , National Institutes of Health, Mammalian Gene Collection (MGC) unpublished (1999)
COMMENT	Contact: Robert Strausberg, Ph.D. Email: cgabs-remail.nih.gov

Tissue Procurement: ATCC
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LNCM1297 row: 3 column: 17
High quality sequence stop: 717.
Location/Qualifiers
1. .955

FEATURES
source
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:4579696"
/clone_lib="NIH_MGC_16"
/tissue_type="retinoblastoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: eye; Vector: pORF7; Site: 1; XhoI; Site: 2; EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: CGCAGCAG(C). Library constructed by Ling Hong in the Laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC Library."
BASE COUNT 268 a 208 c 225 g 254 t
ORIGIN

Query Match 29.1%; Score 29.4; DB 10; Length 955;
Best Local Similarity 66.7%; Pred. No. 1.6e+02;
Matches 42; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

Y 33 ctggatgacgaacgcgggaacccggccagatctacatgcgtcctacgagagca 92
|||
Db 836 CTTCAGATGACTACTCCGAAATCTGCGCGGAAAAAACCGCTTCATGAAGTTGA 895
Y 93 cac 95
|||
Db 896 CAC 898

RESULT 14
BM468546 1000 bp mRNA linear EST 05-FEB-2002
LOCUS
DEFINITION
AGENCOURT_6475521 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:5578160
5', mRNA sequence.
ACCESSION
BM468546
VERSION
BM468546.1 GI:18517588
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
1 (bases 1 to 1000)
NIH-MGC <http://mgc.nci.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgabs-remail.nih.gov
Tissue Procurement: ATCC/DCTD/DTMP
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Plate: LLAM12332 row: m column: 09
High quality sequence stop: 725.
Location/Qualifiers
1. .1000

FEATURES
source
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5578160"

/clone_lib="NIH_MGC_72"
/tissue_type="melanotic melanoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: skin; Vector: pCMV-SPORT6; Site: 1; NotI; Site: 2; SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 2 kb. Library constructed by Life Technologies."
BASE COUNT 276 a 244 c 303 g 174 t 3 others
ORIGIN

Query Match 28.9%; Score 29.2; DB 10; Length 1000;
Best Local Similarity 64.2%; Pred. No. 1.9e+02;
Matches 43; Conservative 0; Mismatches 24; Indels 0; Gaps 0;

Y 22 cccaagctgggctggatgaactcgggaacccggccagatctacatgccttc 81
|||
Db 690 CACAGATGCTGTATGAGAGGACCTCCAGAACTCTGTANACAGACCAGATGAGCTCC 749
Y 82 atgcag 88
|||
Db 750 CAGCAGG 756

RESULT 15
AA264931 313 bp mRNA linear EST 19-APR-2001
LOCUS
DEFINITION
LD08544.5prime LD Drosophila melanogaster embryo Bluescript.
Drosophila melanogaster cDNA clone LD08544 5prime, mRNA sequence.
ACCESSION
AA264931
VERSION
AA264931.1 GI:1900974
KEYWORDS
EST.
SOURCE
fruit fly.
ORGANISM
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE
1 (bases 1 to 313)
Harvey, D., Brokstein, P., Hong, L., Evans-Holm, M., Su, C., Tsang, G.,
Lewis, S. and Rubin, G.M.
BDGP/HHMI Drosophila EST Project
Unpublished (2001)
Contact: Stapleton, M.
BDGP
Lawrence Berkeley National Lab
One Cyclotron Rd, Berkeley, CA 94720, USA
Fax: 510 486 6798
Email: http://www.fruitfly.org/EST_estfruitfly.berkeley.edu
Blast expect value = 1.1E-117 on X03121: Drosophila melanogaster
serendipity (sry) locus DNA sequence
Plate: 85 row: D column: 8
High quality sequence stop: 240.
Location/Qualifiers
1. .313

FEATURES
source
/organism="Drosophila melanogaster"
/db_xref="BDGP_EST:BDCLN007879"
/db_xref="taxon:7227"
/clone="LD08544"
/clone_lib="LD Drosophila melanogaster embryo Bluescript"
/sex="male and female"
/dev_stage="0 to 24 hours mixed stage embryonic"
/lab_host="SOLR"
/note="Organ: embryo; Vector: Bluescript SK; Site: 1; EcoRI; Site: 2; XhoI; Constructed using Stratagene ZAP-cDNA synthesis kit. Oligo dT-primed and directionally cloned at EcoRI and XhoI in Bluescript SK(+/-)"

BASE COUNT 72 a 99 c 82 g 60 t
ORIGIN

Query Match 28.7%; Score 29; DB 9; Length 313;
Best Local Similarity 71.7%; Pred. No. 1.6e+02;
Matches 38; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

Oy 49 cygaaaaccgcgcgccagatctacatgagcttcacatgagagacacactgcta 101
+ ||| ||||| ||| | | | ||||| | ||| |||
Db 235 ctgaagcccgccgcccgtgtccccgcgctttcagagacctctccgacta 287

Search completed: October 3, 2002, 16:15:08
Job time: 16963 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:22:14 : Search time 180.77 Seconds
(without alignments)
137.241 Million cell updates/sec

Title: US-09-826-581-5_COPY_500_600
Perfect score: 101
Sequence: 1 cgtgcccgagagccacat.....atcgagagacacactgtcta 101

Scoring table:
Gapop 10.0 , Gapext 1.0

Searched: 38353 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767056

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: /cgn2_6/ptodata/2/1na/5A.COMB.seq.*
2: /cgn2_6/ptodata/2/1na/5B.COMB.seq.*
3: /cgn2_6/ptodata/2/1na/6A.COMB.seq.*
4: /cgn2_6/ptodata/2/1na/6B.COMB.seq.*
5: /cgn2_6/ptodata/2/1na/PCRUS.COMB.seq.*
6: /cgn2_6/ptodata/2/1na/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Length	DB ID	Description
1	26.4	26.1	44377 2	US-08-804-227C-7 Sequence 7, Appl1
2	26.4	26.1	44377 2	US-08-804-198-1 Sequence 1, Appl1
3	26	25.7	4403765 4	US-09-103-840A-2 Sequence 2, Appl1
4	25.8	25.5	594 4	US-08-441-507-17 Sequence 17, Appl1
5	25.8	25.5	6085 4	US-09-029-603-4 Sequence 4, Appl1
6	25.2	25.0	2980 2	US-08-463-081B-13 Sequence 13, Appl1
7	25.2	25.0	2980 2	US-08-461-379A-13 Sequence 13, Appl1
8	25.2	25.0	2980 2	US-08-462-390B-13 Sequence 13, Appl1
9	25.2	25.0	2980 3	US-08-463-074B-13 Sequence 13, Appl1
10	25.2	25.0	2980 3	US-08-465-585C-13 Sequence 13, Appl1
11	25.2	25.0	2980 3	US-08-652-446-13 Sequence 13, Appl1
12	24.8	24.6	255 4	US-08-980-823-31 Sequence 31, Appl1
13	24.8	24.6	1364 1	US-08-306-691B-50 Sequence 50, Appl1
14	24.8	24.6	1364 5	PCT-US93-06251-65 Sequence 65, Appl1
15	24.8	24.6	1900 4	US-09-608-285A-47 Sequence 47, Appl1
16	24.8	24.6	2294 4	US-09-608-285A-49 Sequence 49, Appl1
17	24.8	24.6	2371 4	US-09-608-285A-46 Sequence 46, Appl1
18	24.8	24.6	2497 4	US-09-608-285A-51 Sequence 51, Appl1
19	24.8	24.6	2693 4	US-09-608-285A-48 Sequence 48, Appl1
20	24.8	24.6	2721 6	5215881-2 Patent No. 5215881
21	24.8	24.6	2762 4	US-09-608-285A-26 Sequence 26, Appl1
22	24.8	24.6	2762 4	US-09-608-285A-52 Sequence 52, Appl1
23	24.8	24.6	2762 4	US-09-240-639-1 Sequence 1, Appl1
24	24.8	24.6	3123 1	US-08-094-889-2 Sequence 2, Appl1
25	24.8	24.6	3123 1	US-07-945-283-1 Sequence 1, Appl1
26	24.6	24.4	30001 1	US-08-125-468-1 Sequence 1, Appl1
27	24.6	24.4	30001 2	US-08-474-933-1 Sequence 1, Appl1

ALIGNMENTS

28	24.4	24.2	2327	4	US-08-868-435-1	Sequence 1, Appl1
29	24.4	24.2	2327	4	US-08-744-231-1	Sequence 1, Appl1
30	24.4	24.2	31571	1	US-08-323-438-1	Sequence 1, Appl1
31	24.4	24.2	68750	3	US-09-335-409-1	Sequence 1, Appl1
32	24.4	24.2	68750	4	US-09-568-102-1	Sequence 1, Appl1
33	24.4	24.2	68750	4	US-09-567-969-1	Sequence 1, Appl1
34	24.4	24.2	68750	4	US-09-568-480-1	Sequence 1, Appl1
35	24.4	24.2	68750	4	US-09-568-486-1	Sequence 1, Appl1
36	24.4	24.2	68750	4	US-09-568-472-1	Sequence 1, Appl1
37	24.4	24.2	71989	4	US-09-443-501A-2	Sequence 2, Appl1
38	24.2	24.0	4695	2	US-08-231-193A-57	Sequence 57, Appl1
39	24.2	24.0	4695	2	US-08-486-273A-57	Sequence 57, Appl1
40	24.2	24.0	4695	3	US-08-940-086A-57	Sequence 57, Appl1
41	24.2	24.0	4695	4	US-08-940-035A-57	Sequence 57, Appl1
42	24.2	23.8	662	4	US-08-441-507-1	Sequence 1, Appl1
43	23.8	23.6	1406	4	US-09-287-097-1	Sequence 1, Appl1
44	23.8	23.6	3048	5	PCT-US95-14418-1	Sequence 1, Appl1
45	23.8	23.6	3048	5	PCT-US95-15327-1	Sequence 1, Appl1

RESULT 1

US-08-804-227C-7
Sequence 7, Application US/08804227C
Patent No. 5876991
GENERAL INFORMATION:
APPLICANT: Dehoff, Bradley S.
APPLICANT: Kuestoss, Stuart A.
APPLICANT: Rostock, Paul R., Jr.
APPLICANT: Sutton, Kimberly L.
TITLE OF INVENTION: POLYKETIDE SYNTHASE GENES
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: THOMAS G. PLANT 1501
STREET: LILLY CORPORATE CENTER
CITY: INDIANAPOLIS
STATE: IN
COUNTRY: USA
ZIP: 46285
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: ASCII(DOS) Text only
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/804,227C
FILING DATE: February 21, 1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Plant, Thomas, G.
REGISTRATION NUMBER: 35,784
REFERENCE/DOCKET NUMBER: X-8231
TELECOMMUNICATION INFORMATION:
TELEPHONE: 317-276-2459
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 44377 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
MOLECULE TYPE: Linear
FEATURE:
NAME/KEY: CDS
LOCATION: 350..14002
FEATURE:
NAME/KEY: CDS
LOCATION: 14046..20036
FEATURE:
NAME/KEY: CDS
LOCATION: 20110..31284
FEATURE:

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; NAME/KEY: CDS
; LOCATION: 31329..36071
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 36155..41830
US-08-804-227C-7

Query Match      26.1%; Score 26.4; DB 2; Length 44377;
Best Local Similarity 69.2%; Pred. No. 17;
Matches 36; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

OY 22 cccaagctggctggtgatgcgaactgcggaaccgcgcgcacatctaca 73
DB 27484 CCGCTGCTGGGAGCGGCTCGAACTGCCGAGTCCGGTGAACCGGATGTACA 27535

RESULT 2
US-08-804-198-1
; Sequence 1, Application US/08804198
; Patent No. 5945320
; GENERAL INFORMATION:
; APPLICANT: Burgett, Stanley G.
; APPLICANT: Kuhstoss, Stuart A.
; APPLICANT: Rao, Nagaraja R.
; APPLICANT: Richardson, Mark A.
; APPLICANT: Rostock, Paul R., Jr.
; TITLE OF INVENTION: PLATENOLIDE SYNTHASE GENE
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PAUL R. CANTRELL, 1138
; STREET: LILLY CORPORATE CENTER
; CITY: INDIANAPOLIS
; STATE: IN
; COUNTRY: USA
; ZIP: 46285
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: Macintosh
; OPERATING SYSTEM: Macintosh 7.0
; SOFTWARE: Microsoft Word 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/804,198
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: CANTRELL, PAUL R.
; REGISTRATION NUMBER: 36,470
; REFERENCE/DOCKET NUMBER: P9113
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 317-276-3885
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 44377 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 350..14002
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 14046..20036
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 20110..31284
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 31329..36071
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 36155..41830
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US-08-804-198-1

Query Match      26.1%; Score 26.4; DB 2; Length 44377;
Best Local Similarity 69.2%; Pred. No. 17;
Matches 36; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

OY 22 cccaagctggctggtgatgcgaactgcggaaccgcgcgcacatctaca 73
DB 27484 CCGCTGCTGGGAGCGGCTCGAACTGCCGAGTCCGGTGAACCGGATGTACA 27535

RESULT 3
US-09-103-840A-2
; Sequence 2, Application US/09103840A
; Patent No. 6294328
; GENERAL INFORMATION:
; APPLICANT: FLEISCHMAN, Robert D.
; APPLICANT: WHITE, Owen R.
; APPLICANT: FRASER, Claire M.
; APPLICANT: VENTER, John C.
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
; TITLE OF INVENTION: TUBERCULOSIS
; FILE REFERENCE: 24366-20007.00
; CURRENT APPLICATION NUMBER: US/09/103,840A
; CURRENT FILING DATE: 1998-06-24
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: Patentln Ver. 2.1
; SEQ ID NO: 2
; LENGTH: 4403765
; TYPE: DNA
; ORGANISM: Mycobacterium tuberculosis
; FEATURE:
; OTHER INFORMATION: CDC 1551
; OTHER INFORMATION: "n" bases at various positions throughout the sequence
; OTHER INFORMATION: represent a, t, c or g
US-09-103-840A-2

Query Match      25.7%; Score 26; DB 4; Length 4403765;
Best Local Similarity 58.8%; Pred. No. 40;
Matches 47; Conservative 0; Mismatches 33; Indels 0; Gaps 0;

OY 6 cccgcagggcccaattcccaagctggctggatgcagacacgcgcgaaccgcgcga 65
DB 2113570 cccgcacgcgltgatccgaacccgcgltacgctgcagcagatcgcagccgcgcaca 2113629

OY 66 gatcatatggcgttcattgc 85
DB 2113630 gaaccgcatggctcagctgc 2113649

RESULT 4
US-08-441-507-17
; Sequence 17, Application US/08441507
; Patent No. 6214358
; GENERAL INFORMATION:
; APPLICANT: Singh, Mohan Bir;
; APPLICANT: Smith, Penelope; and
; APPLICANT: Knox, Robert Bruce
; TITLE OF INVENTION: Protein Allergens of the Species Cynodon
; TITLE OF INVENTION: Dactylon
; NUMBER OF SEQUENCES: 52
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD LLP
; STREET: 28 State Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
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OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: ASCII text
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/441,507
FILING DATE: 15-May-1995
CLASSIFICATION:
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 07/969,875
FILING DATE: 30-October-1992
ATTORNEY/AGENT INFORMATION:
NAME: Mandragouras, Amy E.
REGISTRATION NUMBER: 36,207
REFERENCE/DOCKET NUMBER: IMI-049DV
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 227-7400
TELEFAX: (617) 742-4214
INFORMATION FOR SEQ ID NO: 17:
SEQUENCE CHARACTERISTICS:
LENGTH: 594 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-441-507-17
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Query Match      25.5%; Score 25.8; DB 4; Length 594;
Best Local Similarity 63.9%; Pred. No. 9.7;
Matches 39; Conservative 0; Mismatches 22; Indels 0; Gaps 0;
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QY 8 cgcagagcccaattcccaagctggctggatgacgaactgcggaaccgcgcctcaga 67
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DB 17 CGTTCGGCCGATGCCAGAGAGCGGAGAGCAAGCTGCGCAGCGGCGGAGCTGA 76
QY 68 t 68
DB 77 T 77
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RESULT 5
US-09-029-603-4
Sequence 4, Application US/09029603
Patent No. 6210935
GENERAL INFORMATION:
APPLICANT: Schupp, Thomas
APPLICANT: Engel, Natalie
APPLICANT: Bietenhader, Jurg
APPLICANT: Toupet, Christine
APPLICANT: Pospiech, Andreas
TITLE OF INVENTION: Staurosporin Biosynthesis Gene Clusters
FILE REFERENCE: 4-20555/A/PCF
CURRENT APPLICATION NUMBER: US/09/029,603
CURRENT FILING DATE: 1998-03-20
EARLIER APPLICATION NUMBER: PCT/EP96/03643
NUMBER OF SEQ ID NOS: 11
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 4
LENGTH: 6085
TYPE: DNA
ORGANISM: Streptomyces longisporoflavus
FEATURE:
NAME/KEY: misc_RNA
LOCATION: (378)..(1665)
OTHER INFORMATION: ORF
FEATURE:
NAME/KEY: misc_RNA
LOCATION: (1747)..(2553)
OTHER INFORMATION: ORF
FEATURE:
NAME/KEY: misc_RNA
LOCATION: (2593)..(4011)
OTHER INFORMATION: ORF
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FEATURE:
NAME/KEY: misc_RNA
LOCATION: (4013)..(4999)
OTHER INFORMATION: ORF
FEATURE:
NAME/KEY: misc_RNA
LOCATION: (5071)..(6085)
OTHER INFORMATION: ORF
US-09-029-603-4
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Query Match      25.5%; Score 25.8; DB 4; Length 6085;
Best Local Similarity 56.5%; Pred. No. 17;
Matches 48; Conservative 0; Mismatches 37; Indels 0; Gaps 0;
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QY 15 cccattcccaagctggctggatgacgaactgcggaaccgcgcctcagatctacat 74
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 3243 ccgctggctcaccctcgccagactcgcgacgagcagcagcctggtcacaacat 3302
QY 75 gcgcctcatgacgagacacactgc 99
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 3303 gaacgcagcagcagcagcgtgtcgtgc 3327
```

```
RESULT 6
US-08-463-081B-13/C
Sequence 13, Application US/08463081B
Patent No. 5871960
Patent No. 5871960 5837487
GENERAL INFORMATION:
APPLICANT: Smith, Kendall A. & Beadling, Carol
TITLE OF INVENTION: Nucleic Acids Encoding CR5 Polypeptide,
NUMBER OF SEQUENCES: 35
CORRESPONDENCE ADDRESS:
ADDRESSER: PRETTY, SCHROEDER & POPLANSKI
STREET: 444 South Flower St. - Suite 1900
CITY: Los Angeles
STATE: California
COUNTRY: USA
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0,
SOFTWARE: Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/463,081B
FILING DATE: 5-JUN-1995
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: US 08/104,736
FILING DATE: 10-AUG-1993
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: US 07/796,066
FILING DATE: 20-NOV-91
ATTORNEY/AGENT INFORMATION:
NAME: Viviana Amzel, Ph. D.
REGISTRATION NUMBER: 30,930
REFERENCE/DOCKET NUMBER: P66 38150 (DART-060)
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 622-7700
TELEFAX: (213) 489-4210
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 2980 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 240..1475
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US-08-463-081B-13

Query Match 25.0%: Score 25.2; DB 2; Length 2980;
Best Local Similarity 71.7%: Pred. No. 22;
Matches 33: Conservative 0; Mismatches 13; Indels 0; Gaps 0;

OY 20 ttcccaagctggctggatgacgaactgcggaaccgcgcacca 65
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DB 2119 TTCGAGCGAGGGGTGGGAGCAGACTGCGGAACCTCCCCCAA 2074

RESULT 7

US-08-461-379A-13/C
; Sequence 13, Application US/08461379A
; Patent No. 5871961
; GENERAL INFORMATION:
; APPLICANT: Smith, Kendall A. & Beadling, Carol
; TITLE OF INVENTION: Nucleic Acids Encoding CR5 Polypeptide,
; TITLE OF INVENTION: Vector and Transformed Cell Thereof, and
; NUMBER OF SEQUENCES: 35
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ratner & Prestia
; CITY: Valley Forge
; STATE: Pennsylvania
; COUNTRY: USA
; ZIP: 19482
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0,
; SOFTWARE: Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/461,379A
; FILING DATE: 5-JUNE-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: USSN 08/330,108; 08/104,736
; APPLICATION NUMBER: 6 07/796,066
; FILING DATE: 27-OCT-1994; 10-AUG-1993 & 20-NOV-91
; ATTORNEY/AGENT INFORMATION:
; NAME: Viviana Amzel, Ph. D.
; REGISTRATION NUMBER: 30,930
; REFERENCE/DOCKET NUMBER: DART-070
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (610)470-0700
; TELEFAX: (610)470-0701
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2980 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 240..1475
US-08-461-379A-13

Query Match 25.0%: Score 25.2; DB 2; Length 2980;
Best Local Similarity 71.7%: Pred. No. 22;
Matches 33: Conservative 0; Mismatches 13; Indels 0; Gaps 0;

OY 20 ttcccaagctggctggatgacgaactgcggaaccgcgcacca 65
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DB 2119 TTCGAGCGAGGGGTGGGAGCAGACTGCGGAACCTCCCCCAA 2074

RESULT 8

US-08-462-390B-13/C

; Sequence 13, Application US/08462390B
; Patent No. 5882894
; GENERAL INFORMATION:
; APPLICANT: Smith, K. A. & Beadling, C.
; TITLE OF INVENTION: Nucleic Acids Encoding CR8 Polypeptide, Vector and
; TITLE OF INVENTION: Transformed Cell Thereof, and Expression Thereof
; NUMBER OF SEQUENCES: 35
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ratner & Prestia
; CITY: Valley Forge
; STATE: Pennsylvania
; COUNTRY: USA
; ZIP: 19482
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/462,390B
; FILING DATE: 5-JUNE-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: USSN 08/330,108
; FILING DATE: 27-OCT-1994
; APPLICATION NUMBER: USSN 08/104,736
; FILING DATE: 10-AUG-1993
; APPLICATION NUMBER: USSN 07/796,066
; FILING DATE: 20-NOV-91
; ATTORNEY/AGENT INFORMATION:
; NAME: Viviana Amzel, Ph. D.
; REGISTRATION NUMBER: 30,930
; REFERENCE/DOCKET NUMBER: DART-040
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (610)407-0700
; TELEFAX: (610)407-0701
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2980 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 240..1475
US-08-462-390B-13

Query Match 25.0%: Score 25.2; DB 2; Length 2980;
Best Local Similarity 71.7%: Pred. No. 22;
Matches 33: Conservative 0; Mismatches 13; Indels 0; Gaps 0;

OY 20 ttcccaagctggctggatgacgaactgcggaaccgcgcacca 65
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 2119 TTCGAGCGAGGGGTGGGAGCAGACTGCGGAACCTCCCCCAA 2074

RESULT 9

US-08-463-074B-13/C
; Sequence 13, Application US/08463074B
; Patent No. 6020155
; GENERAL INFORMATION:
; APPLICANT: Smith, Kendall A. & Beadling, Carol
; TITLE OF INVENTION: Nucleic Acids Encoding CR1 Fusion Protein, Vector and
; NUMBER OF SEQUENCES: 35
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PRETTY, SCHROEDER & POPLAWSKI
; CITY: Los Angeles
; STATE: California
; COUNTRY: USA
; ZIP: 90071
444 South Flower St. - Suite 1


```

: TITLE OF INVENTION: Trivalent Synthesis of Oligonucleotides Containing
: NUMBER OF SEQUENCES: 93
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: SCULLY, SCOTT, MURPHY & PRESSER
: STREET: 400 Garden City Plaza
: CITY: Garden City
: STATE: NY
: COUNTRY: USA
: ZIP: 11530
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: OPERATING SYSTEM: IBM PC compatible
: SOFTWARE: Patent Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: PCT/US93/06251
: FILING DATE: 19930630
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: DIGILIO, Frank S.
: REGISTRATION NUMBER: 31,346
: REFERENCE/DOCKET NUMBER: 8586
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 516-742-4343
: TELEFAX: 516-742-4366
: TELEX: 230 901 SANS UR
: INFORMATION FOR SEQ ID NO: 65:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1364 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: PCT-US93-06251-65

Query Match      24.6%; Score 24.8; DB 5; Length 1364;
Best Local Similarity 54.3%; Pred. NO. 25;
Matches 50; Conservative 0; Mismatches 42; Indels 0; Gaps 0;

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DB 529 GGAGACCTGGTGGCCAGACAGTGGATGGGTGGCACCATGGGGGTATCGAGAGCTGCAT 470
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QY 69 ctacatgcgctcatgacgagagacacactgct 100
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DB 469 CTGCTCCAGCTCATGTGGCGGTAGAGCTGCT 438
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RESULT 15
: US-09-608-285A-47/c
: Sequence 47, Application US/09608285A
: Patent No. 6335013
: GENERAL INFORMATION:
: APPLICANT: Ford, John
: APPLICANT: Mulero, Julio
: APPLICANT: Yeung, George
: TITLE OF INVENTION: METHODS AND MATERIALS RELATING TO CD39-LIKE
: FILE REFERENCE: 28110/36570
: CURRENT APPLICATION NUMBER: US/09/608,285A
: CURRENT FILING DATE: 2000-06-30
: PRIOR APPLICATION NUMBER: 09/583,231
: PRIOR FILING DATE: 2000-05-26
: PRIOR APPLICATION NUMBER: 09/557,800
: PRIOR FILING DATE: 2000-04-25
: PRIOR APPLICATION NUMBER: 09/481,238
: PRIOR FILING DATE: 2000-01-11
: PRIOR APPLICATION NUMBER: 09/370,265
: PRIOR FILING DATE: 1999-08-09
: PRIOR APPLICATION NUMBER: PCT/US99/16180
: PRIOR FILING DATE: 1999-07-16
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: PRIOR APPLICATION NUMBER: 09/350,836
: PRIOR FILING DATE: 1999-07-09
: PRIOR APPLICATION NUMBER: 09/273,447
: PRIOR FILING DATE: 1999-03-19
: PRIOR APPLICATION NUMBER: 09/244,444
: PRIOR FILING DATE: 1999-02-04
: PRIOR APPLICATION NUMBER: 09/122,449
: PRIOR FILING DATE: 1998-07-24
: PRIOR APPLICATION NUMBER: 09/118,205
: PRIOR FILING DATE: 1998-07-16
: NUMBER OF SEQ ID NOS: 60
: SOFTWARE: Patentln Ver. 2.0
: SEQ ID NO 47
: LENGTH: 1900
: TYPE: DNA
: ORGANISM: Homo sapiens
: US-09-608-285A-47

Query Match      24.6%; Score 24.8; DB 4; Length 1900;
Best Local Similarity 53.0%; Pred. No. 27;
Matches 53; Conservative 0; Mismatches 47; Indels 0; Gaps 0;

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DB 551 CTGTCTCCAGGGGTCCGACACAGTACTTGGCTGCGATCTCGAAGTCCCCACACACAG 492
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

QY 61 gccagatctacatgcgctcatgacgagagacacactgct 100
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DB 491 GCTGCTCCCTCTCTCCGATCTATGAGGCCACACACAGCT 452
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Job time: 17131 sec

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LVANGVRAAPLMDSKOSKSPVGMILTDFIIVLHRYRSPVQIYEIOHKIETMBREIY
LOGCFKPLVSIISPNDSLFEAVYTLIKRHLRPLVDPSGNVLIHLTKRLLKPLHIF
GSLPRSPFLYRTIODIGICTFRDLAVLETAPLITADIFVDRVSALPYNVECGV
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BASE COUNT 346 a 502 c 462 g 337 t

ORIGIN

Query Match 100.0%; Score 101; DB 6; Length 1647;
Best Local Similarity 100.0%; Pred. No. 1e-20;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1000 GCTCAAGTTCTGCACATCTTGGTCCCTGCTGCCGCCCTCTCTTACCGCAC 1059

OY 61 tatccaagattggcgcacatccgcagactggctg 101
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Db 1060 TATCCAAGATTGGGCATCGCACATTCGACACTGGCTG 1100

RESULT 2
LOCUS AX099776 2109 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 3 from Patent WO0120003.
ACCESSION AX099776
VERSION AX099776.1 GI:13538810
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 2109)
Andersson, L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Rogel-Galliard, C., Iannuccielli, N., Gellin, J., Le Roy, P. and
Chardon, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 3 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
Location/Qualifiers
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YRTIODIGICTFRDLAVLETAPLITADIFVDRVSALPYNVECGVGLYSRDPYH
ILAAQOTYNNHLDMSVGCALRORTLCLEGVISCPHESLGEVIDRIAREQVHRLVYD
E TQHLGLVSLDIQALVLSPPAGIDALGA"

TITLE
VARIANTS OF THE GAMMA CHAIN OF AMPK, DNA SEQUENCES ENCODING THE
SAME, AND USES THEREOF

JOURNAL

INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
Location/Qualifiers

FEATURES

SOURCE

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/db_xref="taxon:9606"
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/note="unnamed protein product"
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ISPNDSLFEAVYTLIKRHLRPLVDPSGNVLIHLTKRLLKPLHIFGSLPRSPFL
YRTIODIGICTFRDLAVLETAPLITADIFVDRVSALPYNVECGVGLYSRDPYH
ILAAQOTYNNHLDMSVGCALRORTLCLEGVISCPHESLGEVIDRIAREQVHRLVYD
E TQHLGLVSLDIQALVLSPPAGIDALGA"

BASE COUNT 458 a 621 c 560 g 470 t

ORIGIN

Query Match 100.0%; Score 101; DB 6; Length 2109;
Best Local Similarity 100.0%; Pred. No. 1e-20;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gctcaagttcctgcacatcttggctcctgcctgcctgcctcctcctcagcagc 60
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Db 900 GCTCAAGTTCTGCACATCTTGGTCCCTGCTGCCGCCCTCTCTTACCGCAC 959

OY 61 tatccaagattggcgcacatccgcagactggctg 101
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Db 960 TATCCAAGATTGGGCATCGCACATTCGACACTGGCTG 1000

RESULT 3
LOCUS AX099802 2115 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 29 from Patent WO0120003.
ACCESSION AX099802
VERSION AX099802.1 GI:13538836
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 2115)
Andersson, L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Rogel-Galliard, C., Iannuccielli, N., Gellin, J., Le Roy, P. and
Chardon, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 29 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
Location/Qualifiers
1..2115
/organism="Homo sapiens"
/db_xref="taxon:9606"
1..1395
/note="unnamed protein product"
/codon_start=1
/protein_id="CAC35801.1"
/db_xref="GI:13538837"
/translation="MSFLEQENSSWPSPAYTSRIRGRRAKALRMTROKSVSEGE
EPGGEGRSPTAESGTLEPATFPKPTPLAADPAVGTPPTGMDCLPSDCTASAG
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RMFOEHCTCDAMATSSKVIPTFEMIKKAFALVANGVRAAPLMDSKOSKSPVGMILT
DFIIVLHRYRSPVQIYEIOHKIETMBREIYLOGCFKPLVSIISPNDSLFEAVYTLI
KRHLRPLVDPSGNVLIHLTKRLLKPLHIFGSLPRSPFLYRTIODIGICTFRDL
AVLETAPLITADIFVDRVSALPYNVECGVGLYSRDPYHILAAQOTYNNHLDMSV
GCALRORTLCLEGVISCPHESLGEVIDRIAREQVHRLVYDETQHLGLVSLDIQAL
VLSPPAGIDALGA"

TITLE
VARIANTS OF THE GAMMA CHAIN OF AMPK, DNA SEQUENCES ENCODING THE
SAME, AND USES THEREOF
Patent: WO 0120003-A 29 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
Location/Qualifiers

FEATURES

SOURCE

1..2115
/organism="Homo sapiens"
/db_xref="taxon:9606"
1..1395
/note="unnamed protein product"
/codon_start=1
/protein_id="CAC35801.1"
/db_xref="GI:13538837"
/translation="MSFLEQENSSWPSPAYTSRIRGRRAKALRMTROKSVSEGE
EPGGEGRSPTAESGTLEPATFPKPTPLAADPAVGTPPTGMDCLPSDCTASAG
SSTIDVEATEPATMECELEGLLEERPALCULSQAFFPKLGMDELRRKPAQAIY
RMFOEHCTCDAMATSSKVIPTFEMIKKAFALVANGVRAAPLMDSKOSKSPVGMILT
DFIIVLHRYRSPVQIYEIOHKIETMBREIYLOGCFKPLVSIISPNDSLFEAVYTLI
KRHLRPLVDPSGNVLIHLTKRLLKPLHIFGSLPRSPFLYRTIODIGICTFRDL
AVLETAPLITADIFVDRVSALPYNVECGVGLYSRDPYHILAAQOTYNNHLDMSV
GCALRORTLCLEGVISCPHESLGEVIDRIAREQVHRLVYDETQHLGLVSLDIQAL
VLSPPAGIDALGA"

BASE COUNT 460 a 622 c 562 g 471 t

ORIGIN

Query Match 100.0%; Score 101; DB 6; Length 2115;
Best Local Similarity 100.0%; Pred. No. 1e-20;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gctcaagttcctgcacatcttggctcctgcctgcctgcctcctcctcagcagc 60
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Db 906 GCTCAAGTTCTGCACATCTTGGTCCCTGCTGCCGCCCTCTCTTACCGCAC 965

OY 61 tatccaagattggcgcacatccgcagactggctg 101
|||||
Db 966 TATCCAAGATTGGGCATCGCACATTCGACACTGGCTG 1006

RESULT 4
LOCUS AF214519 2115 bp mRNA linear PRI 03-JUN-2000
DEFINITION Homo sapiens AMP-activated protein kinase gamma subunit (PRKAG3)
AF214519
ACCESSION AF214519
VERSION AF214519.1 GI:8215681
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE 1 (bases 1 to 2115)
AUTHORS Milan.D., Jeon.J.T., Looft.C., Amarger.V., Robic.A., Thelander.M., Rogel-Galliard.C., Paul.S., Iannuccielli.N., Rask.L., Ronne.H., Lundstrom.K., Reinsch.N., Gellin.J., Kalm.E., Roy.P.L., Chardon.P. and Andersson.L.
JOURNAL MEDLINE 20280150
PUBMED 10818001
TITLE A mutation in PRKAG3 associated with excess glycogen content in pig skeletal muscle
JOURNAL MEDLINE 20280150
PUBMED 10818001
TITLE 2 (bases 1 to 2115)
AUTHORS Milan.D., Jeon.J.T., Looft.C., Amarger.V., Robic.A., Rogel-Galliard.C., Paul.S., Gellin.J., Lundstrom.K., Reinsch.N., Kalm.E., Le Roy.P., Chardon.P. and Andersson.L.
JOURNAL MEDLINE 20280150
PUBMED 10818001
TITLE Direct Submission
JOURNAL MEDLINE 20280150
PUBMED 10818001
TITLE Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish University of Agricultural Sciences, BMC box 597, Uppsala 751 24, Sweden
FEATURES
source location/Qualifiers
1..2115 /organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="2"
/map="2p"
/tissue_type="skeletal muscle"
1..2115 /gene="PRKAG3"
1..1395 /gene="PRKAG3"
/note="AMPK3"
/codon_start=1
/product="AMP-activated protein kinase gamma subunit"
/protein_id="AA173987.1"
/db_xref="GI:8215682"
/translation="MSFLEQNSMSPSPAVTSSERIRGRRAKALRWTRKSYEEG EPPGEGEGRSPPTAESEGLEATPPTKPTPLAODAPAGCTPTGDCIPSPCTSAAG SSTDDVELATERPATFEAMECELEGLERPALCSQAPFPKILMDKELPGAOITM RMDHETCYDMAATSSKLVIFDTMLEIKKAFALVANGVRAAPLMDSKOSFVGLTI TDFILVLRHYRSPVLOIYEI EOHKIEFMREIYLOGCPPLVSI SPNDLFEAVYTLI KNRIHRLPVDPSGVNVLILTHKRLKFLI FGSILRPSFLYRT IODLIGTFEROL AVVLETPA1LTALDLPVDRVSALPVNVECGOVGLSRFVTHLAQOQYTNHLDMSV GEALRPTLCLEGVLSQPHESLGEVIDIRAREQVHRVLVDETHLLGVYSLSDILO ALVSPADIALGA"
BASE COUNT 460 a 622 c 562 g 471 t
ORIGIN
Query Match 100.0%; Score 101; DB 9; Length 2115;
Best Local Similarity 100.0%; Pred. No. 1e-20;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 gctcaagltctgcacacatcttgltccctgctgcgcgcgcctctctctctacgcac 60
|||||
DB 906 GCTCAAGTTCCTGCACATCTTGGTCCCTGCTGCCGCCGCCCTCTCTACCGCAC 965
QY 61 tatccaagattggcgatcgacacatccgcagacttgctg 101
|||||
DB 966 TATCCAGATTGGGCGATCGCACATTCGAGACTTGGCTG 1006
RESULT 5
HSA249977
LOCUS HSA249977 2290 bp mRNA linear PRI 07-APR-2000
DEFINITION Homo sapiens mRNA for AMP-activated protein kinase gamma 3 subunit (AMPK gamma 3 gene).
ACCESSION AJ249977
VERSION AJ249977.1 GI:6688200
KEYWORDS AMP-activated protein kinase; AMPK gamma 3 gene; gamma 3 subunit.
SOURCE human
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 2290)
AUTHORS Cheung.P.C., Salt.I.P., Davies.S.P., Hardie.D.C. and Carling.D.
TITLE Characterization of AMP-activated protein kinase gamma-subunit isoforms and their role in AMP binding
JOURNAL MEDLINE 20164049
PUBMED 10818001
TITLE 2 (bases 1 to 2290)
AUTHORS Carling.D.
JOURNAL MEDLINE 20164049
PUBMED 10818001
TITLE Direct Submission
JOURNAL MEDLINE 20164049
PUBMED 10818001
TITLE Submitted (12-OCT-1999) Carling D., Cellular Stress Group, MRC Clinical Sciences Centre, Hammersmith Hospital, DuCane Road, London, W12 0NN, UNITED KINGDOM
FEATURES
source location/Qualifiers
1..2290 /organism="Homo sapiens"
/db_xref="taxon:9606"
22..1500 /gene="AMPK gamma 3"
22..1500 /gene="AMPK gamma 3"
/function="AMP-activated protein kinase regulatory subunit"
/codon_start=1
/evidence="experimental"
/product="AMP-activated protein kinase gamma 3 subunit"
/protein_id="CAB5117.1"
/db_xref="GI:6688201"
/translation="MEGLHALRRTPSMSSLGSEHOENMSFLEQNSMSPSPAVTS SSERIRGRRAKALRWTRKSYEEGEGEGRSPPTAESEGLEATPPTKPTPLAODAPAGCTPTGDCIPSPCTSAAG SSTDDVELATERPATFEAMECELEGLERPALCSQAPFPKILMDKELPGAOITM RMDHETCYDMAATSSKLVIFDTMLEIKKAFALVANGVRAAPLMDSKOSFVGLTI TDFILVLRHYRSPVLOIYEI EOHKIEFMREIYLOGCPPLVSI SPNDLFEAVYTLI KNRIHRLPVDPSGVNVLILTHKRLKFLI FGSILRPSFLYRT IODLIGTFEROL AVVLETPA1LTALDLPVDRVSALPVNVECGOV GLSRFVTHLAQOQYTNHLDMSV GEALRPTLCLEGVLSQPHESLGEVIDIRAREQVHRVLVDETHLLGVYSLSDILOALVSPADIALGA"
BASE COUNT 501 a 674 c 617 g 498 t
ORIGIN
Query Match 100.0%; Score 101; DB 9; Length 2290;
Best Local Similarity 100.0%; Pred. No. 1e-20;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 gctcaagltctgcacacatcttgltccctgctgcgcgcgcctctctctctacgcac 60
|||||
DB 1002 GCTCAAGTTCCTGCACATCTTGGTCCCTGCTGCCGCCGCCCTCTCTACCGCAC 1061
QY 61 tatccaagattggcgatcgacacatccgcagacttgctg 101
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DB 1062 TATCCAGATTGGGCGATCGCACATTCGAGACTTGGCTG 1102
RESULT 6
AX099774
LOCUS AX099774 1867 bp DNA linear PAT 02-APR-2001
DEFINITION Sequence 1 from Patent W00120003.
ACCESSION AX099774
VERSION AX099774.1 GI:13538808
KEYWORDS pig.
SOURCE Sus scrofa
ORGANISM Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Sus.
REFERENCE 1 (bases 1 to 1867)
AUTHORS Andersson.L., Looft.C., Kalm.E., Milan.D., Robic.A., Rogel-Galliard.C., Iannuccielli.N., Gellin.J., Le Roy.P. and Chardon.P.
JOURNAL MEDLINE 20164049
PUBMED 10818001
TITLE Variants of the gamma chain of ampk, dna sequences encoding the same, and uses thereof
PATENT: WO 0120003-A 1 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ; ANDERSSON, Leif (SE) ; LOOFT, Christian (DE) ; KALM, Ernst (DE)

FEATURES
source
1..1867
/organism="Sus scrofa"
/db_xref="taxon:9823"
472..1389
CDS
/note="unnamed protein product"
/codon_start=1
/protein_id="CAC35798.1"
/db_xref="GI:13538809"
/translation="MHFQEHCTIDAMATSSKLVIFDTMLEIKKAFALVANGVRAAP
LSDSKKQSFVGMVLTDFILVLRYSRPLVOIYEIEHKIETWREILQGCFFPLVS
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YRTIIDLIGITFRLAVLEFAPILTLADITFVDRVSALPVNENGVGVGLYSRFDV
HLAAGOTYNHLDNMVGFALRORTLCIEGLVSCQPHETLGEVIDRIYRQVIRLVLYDE
TQHLGVSLSLDILOALVLSFAGIDALGA"

BASE COUNT 380 a 583 c 529 g 375 t
ORIGIN

Query Match 92.7% Score 93.6: DB 6: Length 1867;
Best Local Similarity 96.0% Pred. No. 1.9e-18;
Matches 96: Conservative 0: Mismatches 4: Indels 0: Gaps 0:

QY 2 ctcaagttccgcacacatttggtccctgctgcgcgcgcgcctcctcctaccgcact 61
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DB 901 CTCAGTTCCTGCACATCTTGGCACCTGCTGCCGCCCTCCTCTTACCGCAC 960
|||||
QY 62 atccaagattgggcacatccgcacatccgcagactggctg 101
|||||
DB 961 ATCCAAGATTGGGCATCGCACATTCCGACACTTGGCGC 1000
|||||

RESULT 7
AF214520 1873 bp mRNA linear MAM 03-JUN-2000
LOCUS
DEFINITION Sus scrofa AMP-activated protein kinase gamma subunit (PRKAG3)
ACCESSION AF214520
VERSION AF214520
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
AUTHORS
1 (bases 1 to 1873)
Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Thelander,M.,
Rogel-Gallard,C., Paul,S., Jannuccelli,N., Rask,L., Ronne,H.,
Lundstrom,K., Reinsch,N., Gellin,J., Kaim,E., Roy,P.L., Chardon,P.
and Andersson,L.
A mutation in PRKAG3 associated with excess glycogen content in pig
skeletal muscle
Science 288 (5469), 1248-1251 (2000)
JOURNAL
MEDLINE
PUBMED
REFERENCE
AUTHORS
2 (bases 1 to 1873)
Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A.,
Rogel-Gallard,C., Paul,S., Gellin,J., Lundstrom,K., Reinsch,N.,
Kaim,E., Le Roy,P., Chardon,P. and Andersson,L.
Direct Submission
TITLE
JOURNAL
Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish
University of Agricultural Sciences, BMC Box 597, Uppsala 751 24,
Sweden
FEATURES
source
1..1873
/organism="Sus scrofa"
/db_xref="taxon:9823"
/chromosome="15"
/map="15q"
/tissue-type="skeletal muscle"
1..1873
/gene="PRKAG3"
1..1395
/gene="PRKAG3"

/note="AMPK3"
/codon_start=1
/product="AMP-activated protein kinase gamma subunit"
/protein_id="AA073988.1"
/db_xref="GI:8215684"
/translation="MSFLEGGESRSPWSPRAVTTSSERSHGDCNKKASRMTROEDVEEG
GPGPREGPOSRPVASTGOEATFPKATPLAQAAPLAEVONPPTERRDILSDCAASAS
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DFILVLRYSRPLVOIYEIEHKIETWREILQGCFFPLVSISPNDLSFEAYVALIKNR
ILHRLPYLDVPSGAVLHILTHKRLKFLHIGTLPRPSFLYRTIIDLIGITFRLAVLE
FAPILTLADITFVDRVSALPVNENGVGVGLYSRFDVHLAAGOTYNHLDNMVGFAL
RORTLCIEGLVSCQPHETLGEVIDRIYRQVIRLVLYDETHLGVLSLDILOALVLSF
AGIDALGA"

BASE COUNT 382 a 580 c 535 g 376 t
ORIGIN

Query Match 92.7% Score 93.6: DB 4: Length 1873;
Best Local Similarity 96.0% Pred. No. 1.9e-18;
Matches 96: Conservative 0: Mismatches 4: Indels 0: Gaps 0:

QY 2 ctcaagttccgcacacatttggtccctgctgcgcgcgcgcctcctcctaccgcact 61
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DB 907 CTCAGTTCCTGCACATCTTGGCACCTGCTGCCGCCCTCCTCTTACCGCAC 966
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QY 62 atccaagattgggcacatccgcacatccgcagactggctg 101
|||||
DB 967 ATCCAAGATTGGGCATCGCACATTCCGACACTTGGCGC 1006
|||||

RESULT 8
AX099800 1873 bp DNA linear PAT 02-APR-2001
LOCUS
DEFINITION Sequence 27 from Patent WO0120003.
ACCESSION AX099800
VERSION AX099800
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
AUTHORS
1 (bases 1 to 1873)
Andersson,L., Looft,C., Kaim,E., Milan,D., Robic,A.,
Rogel-Gallard,C., Jannuccelli,N., Gellin,J., Le Roy,P. and
Chardon,P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 27 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kaim, Ernst (DE)
TITLE
JOURNAL
Location/Qualifiers
1..1873
/organism="Sus scrofa"
/db_xref="taxon:9823"
1..1395
/note="unnamed protein product"
/codon_start=1
/protein_id="CAC35800.1"
/db_xref="GI:13538835"
/translation="MSFLEGGESRSPWSPRAVTTSSERSHGDCNKKASRMTROEDVEEG
GPGPREGPOSRPVASTGOEATFPKATPLAQAAPLAEVONPPTERRDILSDCAASAS
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HFMOEHCTIDAMATSSKLVIFDTMLEIKKAFALVANGVRAAPLSDSKKQSFVGMVLT
DFILVLRYSRPLVOIYEIEHKIETWREILQGCFFPLVSISPNDLSFEAYVALIKNR
ILHRLPYLDVPSGAVLHILTHKRLKFLHIGTLPRPSFLYRTIIDLIGITFRLAVLE
FAPILTLADITFVDRVSALPVNENGVGVGLYSRFDVHLAAGOTYNHLDNMVGFAL
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AGIDALGA"

FEATURES
source
1..1873
/organism="Sus scrofa"
/db_xref="taxon:9823"
1..1395
/note="unnamed protein product"
/codon_start=1
/protein_id="CAC35800.1"
/db_xref="GI:13538835"
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GPGPREGPOSRPVASTGOEATFPKATPLAQAAPLAEVONPPTERRDILSDCAASAS
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HFMOEHCTIDAMATSSKLVIFDTMLEIKKAFALVANGVRAAPLSDSKKQSFVGMVLT
DFILVLRYSRPLVOIYEIEHKIETWREILQGCFFPLVSISPNDLSFEAYVALIKNR
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FAPILTLADITFVDRVSALPVNENGVGVGLYSRFDVHLAAGOTYNHLDNMVGFAL
RORTLCIEGLVSCQPHETLGEVIDRIYRQVIRLVLYDETHLGVLSLDILOALVLSF
AGIDALGA"

BASE COUNT 382 a 580 c 535 g 376 t
ORIGIN

Query Match 92.7% Score 93.6; DB 6; Length 1873;
Best Local Similarity 96.0% Pred. No. 1.9e-18;
Matches 96; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ctcaagttccgcacatcttggtctccctgctgcgcgcgcctctctctctaccgact 61
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Db 907 CTCAGTTCCTCGCACATCTTTGGCACCTCTGCTGCCCGCCCTCTCTCTACCGCACC 966
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QY 62 atccaagattggcgcacatctccgagacttgctg 101
|||||
Db 967 ATCCAGATTGGGCATCGCACATTCGAGACTTGGCCG 1006
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RESULT 9
AX099804 2022 bp DNA linear PAT 02-APR-2001
LOCUS AX099804
DEFINITION Sequence 31 from Patent WO0120003.
ACCESSION AX099804
VERSION AX099804.1 GI:13538838
KEYWORDS
SOURCE
ORGANISM
Sus scrofa
Pig.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
Andersson L., Looft, C., Kalm, E., Milan, D., Robic, A.,
Rogel-Galliard, C., Iannucci, N., Geillin, J., Le Roy, P. and
Charidon, P.
Variants of the gamma chain of ampk, dna sequences encoding the
same, and uses thereof
Patent: WO 0120003-A 31 22-MAR-2001;
INSTITUT NATIONAL DE LA RECHERCHE AGRONOMIQUE (INRA) (FR) ;
Andersson, Leif (SE) ; Looft, Christian (DE) ; Kalm, Ernst (DE)
Location/Qualifiers
1. .2022
/organism="Sus scrofa"
/db_xref="taxon:9823"
BASE COUNT 412 a 623 c 593 g 394 t
ORIGIN

Query Match 92.7% Score 93.6; DB 6; Length 2022;
Best Local Similarity 96.0% Pred. No. 1.9e-18;
Matches 96; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ctcaagttccgcacatcttggtctccctgctgcgcgcgcctctctctaccgact 61
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Db 1057 CTCAGTTCCTCGCACATCTTTGGCACCTCTGCTGCCCGCCCTCTCTACCGCACC 1116
|||||

QY 62 atccaagattggcgcacatctccgagacttgctg 101
|||||
Db 1117 ATCCAGATTGGGCATCGCACATTCGAGACTTGGCCG 1156
|||||

RESULT 10
AX281580 1722 bp DNA linear PAT 03-NOV-2001
LOCUS AX281580
DEFINITION Sequence 3 from Patent WO0177305.
ACCESSION AX281580
VERSION AX281580.1 GI:16608831
KEYWORDS
SOURCE
ORGANISM
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (sites)
Andersson, L., Luthman, H. and Marklund, S.
TITLE
Variants of the human amp-activated protein kinase gamma 3 subunit
JOURNAL
Patient: WO 0177305-A 3 18-OCT-2001;
Aresis AB (SE)
Location/Qualifiers
1. .1722
/organism="Homo sapiens"

BASE COUNT 321 a 504 c 534 g 363 t
ORIGIN /db_xref="taxon:9606"

Query Match 78.4% Score 79.2; DB 6; Length 1722;
Best Local Similarity 91.3% Pred. No. 4.9e-14;
Matches 84; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 10 cctgcacatcttggtctccctgctgcgcgcgcctctctctaccgactccaga 69
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Db 1510 CCATCCTACACAGAGGTTCCCTGCTGCCCGCCCTCTCTCTACCGACTATCCACA 1569
|||||

QY 70 ttgggcgcgcacatctccgagacttgctg 101
|||||
Db 1570 TTTGGGCATCGCACATTCGAGACTTGGCTG 1601
|||||

RESULT 11
AC027416/c 152129 bp DNA linear HTG 07-JUN-2000
LOCUS AC027416
DEFINITION Homo sapiens clone RP11-504G11, WORKING DRAFT SEQUENCE, 32
unordered pieces.
ACCESSION AC027416
VERSION AC027416.2 GI:8317289
KEYWORDS
SOURCE
ORGANISM
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 152129)
Birren, B., Linton, L., Nusbaum, C. and Lander, E.
TITLE
JOURNAL
Homo sapiens, clone RP11-504G11
REFERENCE
2 (bases 1 to 152129)
1 (bases 1 to 152129)
Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
Anderson, S., Baldwin, J., Barna, N., Bastien, V., Bede, F.,
Boguslavsky, L., Boukhalter, B., Brown, A., Burkett, G.,
Campopiano, A., Castle, A., Choepel, Y., Colangelo, M., Collins, S.,
Collamore, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S.,
Dodge, S., Domino, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D.,
Galagan, J., Gardyna, S., Glade, S., Goyette, M., Graham, L.,
Grand-Pierre, N., Grant, G., Hags, B., Heatford, A., Horton, L.,
Howard, J.C., Iliev, I., Johnson, R., Jones, C., Kann, L., Karatas, A.,
Klein, J., Lacombe, K., Lamazares, R., Landers, T., Lehoczy, J.,
Levine, R., Lieu, C., Liu, G., Locke, K., MacDonald, P., Marquis, N.,
McCarthy, M., McKean, P., McGurk, A., McKernan, K., McPheters, R.,
Meldrum, J., Menus, L., Mihova, T., Miranda, C., Miñana, V., Morrow, J.,
Murphy, T., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell, P.,
O'Neill, D., Olivari, T.M., Oliver, J., Peterson, K., Pierre, N.,
Pisanil, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D.,
Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B.,
Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talmas, J.,
Testfay, S., Theodore, J., Tirrell, A., Travers, M., Triggillo, J.,
Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J.,
Young, G., Zainoun, J., Zimmer, A. and Zody, M.
Submitted (30-MAR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 7, 2000 this sequence version replaced gi:7342115.
COMMENT
TITLE
JOURNAL
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/BW/RepeatMasker.html

Center: Whitehead Institute / MIT Center for Genome Research
Center code: W18R
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu

Project Information
Center project name: L7458
Center clone name: 504_G.11

Summary Statistics
Sequencing vector: M13; M77815; 100% of reads

Query Match	78.4%;	Score 79.2;	DB 2;	Length 196554;
Best Local Similarity	91.3%;	Pred. No. 5.2e-14;		
Matches 84;	Conservative	0;	Mismatches 8;	Indels 0;
				Gaps 0;

10 ccgcacaccttgggtccctgctgcccgccctccctcctctacgcgaactatccaaga 65

59610 CCATCCTAACCAAGGTTCCCTGCTGCCCGGCCCTCCTTCTCTACCGCACTATCCAAGA 59551

70 cttgggcatcggcacaatccgaagacttgctg 101
|||||
59550 tttgggcacatcgacacattccgagacttgctg 59519

[illegible]

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
1 (bases 1 to 206854)
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)

MEDLINE	99063792
REFERENCE	2 (bases 1 to 206654)
AUTHORS	Harris, A. and Colton, M.
TITLE	The Sequence of Homo sapiens BAC clone RP11-459119

JOURNAL
REFERENCE
AUTHORS
TITLE
Submitted (08-SEP-1999) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis
MO 63108, USA
4 (bases 1 to 206854)

AUTHORS Waterston, R. H.
TITLE Direct Submission
Submitted (08-NOV-2001) Genome Sequencing Center, Washington
JOURNAL University School of Medicine, 4444 Forest Park Parkway, St. Louis

MO 63108, USA
5 (bases 1 to 206854)
Waterson, R.H.
Direct Submission
Submitted (03-JAN-2002) Genome Sequencing Center, Washington

University School of Medicine, 4444 Forest Park Parkway, St. Louis
MO 63108, USA
6 (bases 1 to 206854)

AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (09-JAN-2002) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
COMMENT On Nov 8, 2001 this sequence version replaced g1:13433203.

----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc>
Contact: saplens@wustl.edu
----- Summary Statistics
Center project name: H_NH0459119

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30): an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McClellerson, Department of Genetics, Washington University, St. Louis, MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION: The Rpci-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Moon, P.-Y., Zhao, B., Frengen, E., Tanno, M., Catanesi, J. J. and de Jong, P. J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is Rp11-1077K22; the clone sequenced to the right is Rp11-64705. Actual start of this clone is at base position 1 of Rp11-459119; actual end is at base position 206854 of Rp11-459119.

Dataform A0799810 and AC073128 was used to finish this clone, and
 AC009974. Polymorphisms have been identified between AC073128 and
 AC009974. A single plasmid region exists between 58812-58903.
 A single plasmid region exists between 184390-185163.
 The difference between the two plasmids is approximately 1700 bps.
 PCR suggests that approximately 1700 bps are missing.

FEATURES

Source

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Db 165190	CCATCTTAACGACAGGGTTCCTCCTGTCCCCGGCGCTTCCTCTAACGCCACTATTCCAAGA	165131											
Oy	70	tttggcatcgccacattccgcgaacttgctg	101										
Db 165130	TTTGGCATCGGCACATTCCGAACTTGCTG	165099											
RESULT	14												
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LOCUS	AF214521	Sus scrofa	AMPK gamma subunit (PRKAG3)	gene, complete cds.									
DEFINITION	AF214521												
ACCESSION	AF214521.1	GI:8215685											
VERSION													
KEYWORDS	.												
SOURCE	pig.												
ORGANISM	Sus scrofa												
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;												
AUTHORS	Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.												
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	Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A., Thelander,M.,												
	Rogel-Gallard,C., Paul,S., Tannucci,L.N., Rask,L., Ronne,H.,												
	Lundstrom,K., Reinsch,N., Gellin,J., Kalm,E., Roy,P.L., Chardon,P.												
	and Andersson,L.												
TITLE	A mutation in PRKAG3 associated with excess glycogen content in pig												
JOURNAL	skeletal muscle												
MEDLINE	Science 288 (5469), 1248-1251 (2000)												
PUBMED	20280150												
REFERENCES	10818001												
AUTHORS	2 (bases 1 to 5888)												
	Milan,D., Jeon,J.T., Looft,C., Amarger,V., Robic,A.,												
	Rogel-Gallard,C., Paul,S., Gellin,J., Lundstrom,K.,												
	Kalm,E., Le Roy,P., Chardon,P. and Andersson,L.												
TITLE	Direct Submission												
JOURNAL	Submitted (10-DEC-1999) Dept Animal Breeding and Genetics, Swedish												
	University of Agricultural Sciences, BMC Box 597, Uppsala 751 24,												
	Sweden												
FEATURES													
source	Location/Qualifiers												
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Search completed: October 3, 2002, 14:50:23
 Job time: 12143 sec

Query Match 75.2%; Score 76; DB 4; Length 5888;

Best Local Similarity 85.0%; Pred. No. 4.7e-13;

Matches 85; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

OY 2 ctcaagtlcctgacatcttgllccctgcccgcctcctcctcctcact 61

DB 3265 CACGCTCCCAACCAACGACACCTGCTGCCGCCCTCTTCTTACCGCAC 3324

OY 62 atccaagattggcagcagcattccgagacttgctg 101

DB 3325 ATCCAGATTGGCATCGCACATTCCGACACTTGCCG 3364

RESULT 15

AF336381/C

LOCUS AF336381 227724 bp DNA linear HTG 02-APR-2001

DEFINITION Mus musculus chromosome 1 clone PAC510; PAC457, *** SEQUENCING IN

PROGRESS ***, 3 unordered pieces.

ACCESSION AF336381 GI:13507298

VERSION HTG; HTGS_PHASE1.

KEYWORDS house mouse.

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

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REFERENCE

BASE COUNT 57663 a 55423 c 56238 g 58199 t 201 others

ORIGIN

Query Match 70.3%; Score 71; DB 2; Length 227724;

Best Local Similarity 93.7%; Pred. No. 1.7e-11;

Matches 74; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 23 ggttcctgctgcccgcctcctcctcctcactatccagattggcagc 82

DB 5531 GGTGCTCTGTTGCCCGGCTCTCTGCGCACTATCCAGACTTGCGGC 5472

OY 83 acattccgagacttgctg 101

DB 5471 ACATTCCGAGATTGCGTG 5453

FEATURES

source

1. 227724

/organism="Mus musculus"

/db_xref="taxon:10090"

/chromosome="1"

/clone="PAC510; PAC457"

Submitted (17-JAN-2001) Genome Analysis, Institute of Molecular
 Biotechnology, Beutenbergstr. 11, Jena 07745, Germany
 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 3 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

* 1 17869: contig of 17869 bp in length

* 17870 17969: gap of unknown length

* 32746: contig of 14777 bp in length

* 32747 32846: gap of unknown length

* 32847 227724: contig of 194878 bp in length.

Location/Qualifiers

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:30:46 ; Search time 719.93 Seconds
(without alignments)
240.868 Million cell updates/sec

Title: US-09-826-581-5_COPY_1000_1100
Perfect score: 101
Sequence: 1 gctcaagttcttcgacatct.....cacatccgagacttgcctg 101

Scoring table:
IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1736436 seqs, 858457221 residues
Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: N.Geneseq_032802.*
2: /SIDSL/gcgdata/geneseq/geneseqn-emb1/NA1980.DAT.*
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24: /SIDSL/gcgdata/geneseq/geneseqn-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	101	100.0	547	22	ABAO8485
2	101	100.0	1647	22	AAH43685
3	101	100.0	2109	22	AAH43685
4	101	100.0	2115	22	AAH43685
5	93.6	92.7	1867	22	AAH43685
6	93.6	92.7	1873	22	AAH43685
7	93.6	92.7	2022	22	AAH43685
8	79.2	78.4	1722	22	AAH43685
9	30.8	30.5	1435	20	AAH43685

10	30.8	30.5	1467	23	AAH43685	DNA encoding novel
11	30.8	30.5	2223	22	AAH43685	Human CDNA sequenc
12	29.6	29.3	566	22	AAH43685	Human CDNA sequenc
13	29.6	29.3	1326	21	AAH43685	Human CDNA sequenc
14	29.6	29.3	2289	23	AAH43685	Human CDNA sequenc
15	29.6	29.3	4363	23	AAH43685	Human CDNA sequenc
16	29.4	29.1	200	24	AAH43685	Human CDNA sequenc
17	29.4	29.1	1259	14	AAH43685	Human CDNA sequenc
18	29.4	29.1	1260	20	AAH43685	Human CDNA sequenc
19	29.4	29.1	1260	20	AAH43685	Human CDNA sequenc
20	29.4	29.1	1260	24	AAH43685	Human CDNA sequenc
21	29.4	29.1	5007	23	AAH43685	Human CDNA sequenc
22	29.4	29.1	12403	23	AAH43685	Human CDNA sequenc
23	29.4	29.1	4413	23	AAH43685	Human CDNA sequenc
24	29.4	29.1	6765	23	AAH43685	Human CDNA sequenc
25	29.4	29.1	7461	23	AAH43685	Human CDNA sequenc
26	28.8	28.5	398	22	AAH43685	Human CDNA sequenc
27	28.8	28.5	9979	22	AAH43685	Human CDNA sequenc
28	28.8	28.5	13673	22	AAH43685	Human CDNA sequenc
29	28.8	28.5	13673	22	AAH43685	Human CDNA sequenc
30	28.4	28.1	1682	21	AAH43685	Human CDNA sequenc
31	28.2	27.9	635	22	AAH43685	Human CDNA sequenc
32	28.2	27.9	2378	23	AAH43685	Human CDNA sequenc
33	28.2	27.9	2530	21	AAH43685	Human CDNA sequenc
34	28.2	27.9	2588	23	AAH43685	Human CDNA sequenc
35	28.2	27.9	2857	22	AAH43685	Human CDNA sequenc
36	28.2	27.9	3238	23	AAH43685	Human CDNA sequenc
37	28.2	27.9	4378	23	AAH43685	Human CDNA sequenc
38	28.2	27.9	4732	23	AAH43685	Human CDNA sequenc
39	28.2	27.7	301	23	AAH43685	Human CDNA sequenc
40	28.2	27.7	737	22	AAH43685	Human CDNA sequenc
41	28.2	27.7	3125	22	AAH43685	Human CDNA sequenc
42	28.2	27.7	3156	19	AAH43685	Human CDNA sequenc
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45	28.2	27.7	12043	24	AAH43685	Human CDNA sequenc

ALIGNMENTS

RESULT 1	ABAO8485	standard; CDNA: 547 BP.
ID	ABAO8485	11-JAN-2002 (first entry)
AC	ABAO8485	
DT	11-JAN-2002	
XX		
DE	Human AMP-activated protein kinase subunit homologue CDNA, SEQ ID NO:261.	
XX		
KW	Human; cytokine; cell proliferation; cell differentiation; growth factor;	
KW	hematopoiesis regulation; tissue growth; immunomodulator; activin;	
KW	inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;	
KW	proliferation; metastasis; cancer; tumor; hematopoietic disorder;	
KW	myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;	
KW	chronic inflammatory condition; proliferative retinopathy;	
KW	atherosclerosis; coronary heart disease; arterial ischemia;	
KW	bone disorder; osteoporosis; vascular growth disorder;	
KW	tissue regeneration; wound healing; infection; immune disorder;	
KW	cell culture; drug screening; gene therapy; antiinflammatory;	
KW	antisthmatic; antiarthritic; hemostatic; antiarteriosclerotic;	
KW	cytostatic; osteopathic; vasotropic; cardiant; virucide; antibacterial;	
KW	antifungal; vulnery; antitumor; ss.	
XX		
OS	Homo sapiens.	
PN	WO200157188-A2.	
XX		
PD	09-AUG-2001.	
XX		
PF	05-FEB-2001; 2001WO-US03800.	
XX		

PR 03-FEB-2000: 2000US-0496914.
 PR 27-APR-2000: 2000US-0560875.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Tang YT, Liu C, Drmanac RT;
 XX
 DR WPI: 2001-457740/49.
 DK P-PSDB: ABH1241.
 XX
 PT Human proteins and DNA encoding sequences useful for preventing,
 PT treating or ameliorating a medical condition in a mammalian subject
 PT e.g. arthritis and cancer -
 XX
 PS Claim 1: Page 429: 1963pp: English.
 XX
 CC Sequences ABH10981-ABH12330 represent 1350 novel human polypeptides, and
 CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The
 CC invention also relates to vectors and recombinant host cells comprising a
 CC nucleotide of the invention, methods of producing the novel polypeptides,
 CC antibodies against the polypeptides, methods of detecting the nucleotides
 CC or polypeptides in a sample, and methods of identifying compounds which
 CC bind to polypeptides of the invention. Although novel, many of the
 CC polypeptides of the invention have homology to known proteins, thereby
 CC giving an insight into their probable biological activities, and hence
 CC potential therapeutic applications. The polypeptides of the invention may
 CC have various activities, including cytokine, cell proliferation or cell
 CC differentiation activities; stem cell growth factor activity;
 CC haematopoiesis regulatory activity; tissue growth factor activity;
 CC immunomodulatory activity; activin- or inhibin-related activities;
 CC chemotactic or chemokinetic activities; haemostatic, thrombotic or
 CC thrombolytic activities; receptor or ligand activities; or may be
 CC involved in oncogenesis, cancer cell proliferation or metastasis.
 CC Depending on their biological activities, polypeptides and nucleotides of
 CC the invention are useful for preventing, treating or ameliorating medical
 CC conditions, e.g. by protein or gene therapy. Such conditions include
 CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell
 CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),
 CC proliferative retinopathy, atherosclerosis, coronary heart disease,
 CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal
 CC vascular growth. Polypeptides involved with tissue regeneration and
 CC repair (or nucleic acids encoding them) may be used to promote wound
 CC healing (e.g., of burns, incisions and ulcers), while those with
 CC immunomodulatory activities may be used in the treatment of viral,
 CC bacterial and fungal infections in addition to immune disorders.
 CC Polypeptides with growth factor activity may be used in cell cultures to
 CC promote cell growth. For example, such polypeptides may be used to
 CC manipulate stem cells in culture to give rise to neuroepithelial cells
 CC that can be used to augment or replace cells damaged by illness,
 CC autoimmune disease or accidental damage. The polypeptides and nucleotides
 CC may also be used in the diagnosis of the above conditions, and in drug
 CC screening techniques. The present sequence represents a cDNA encoding a
 CC novel human polypeptide of the invention.
 XX
 SQ Sequence 547 BP: 112 A: 172 C: 133 G: 130 T: 0 other;

Query Match 100.0%; Score 101; DB 22; Length 547;
 Best Local Similarity 100.0%; Pred. No. 7.9e-23;
 Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gctcaagttcctgcacatcttggtccctgcgcgcgcgcctctctctacgcac 60
 |||||||
 DB 263 gctcaagttcctgcacatcttggtccctgcgcgcgcgcctctctctacgcac 322
 |||||||
 OY 61 tatccaagatttggcgcgcgcacatctcgagactgctg 101
 |||||||
 DB 323 tatccaagatttggcgcgcgcacatctcgagactgctg 363
 |||||||

RESULT 2
 AAH43685
 ID AAH43685 standard; cDNA: 1647 BP.

XX
 AC AAH43685;
 XX
 DT 21-JAN-2002 (first entry)
 XX
 DE PRKAG3 cDNA.
 XX
 KM Human; AMP-activated protein kinase gamma 3 subunit; PRKAG3; variant;
 KM metabolic disease; diabetes; obesity; substitution; ss.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT 20..1489
 FT /*tag= a
 FT /product= "PRKAG3"
 FT 230
 FT /*tag= b
 FT /label= "C230C"
 FT /note= "Causes P71A"
 FT 559
 FT /*tag= c
 FT /label= "T559C"
 FT /note= "Silent variation"
 FT 1037
 FT /*tag= d
 FT /label= "C1037T"
 FT /note= "Causes R340W"
 XX
 PN W0200177305-A2.
 XX
 PD 18-OCT-2001.
 XX
 PF 06-APR-2001; 2001WO-SE00765.
 XX
 PR 07-APR-2000; 2000US-195665P.
 XX
 PA (AREX-) AREXIS AB.
 XX
 PI Andersson L, Luthman H, Marklund S;
 XX
 DR WPI: 2001-657170/75.
 DR P-PSDB: Q0B47679.
 XX
 PT New variants of human AMP-activated protein kinase gamma3 subunit
 PT associated with a metabolic disease e.g. diabetes or obesity and method
 PT for determining a risk estimate of diseases in subject by detecting the
 PT variant -
 XX
 PS Disclosure; Fig 5: 25pp; English.
 XX
 CC This sequence represents the full length cDNA encoding the human
 CC AMP-activated protein kinase gamma 3 subunit (PRKAG3). Detecting
 CC the presence of the PRKAG3 DNA, or a variant, is useful in determining
 CC a risk estimate of a metabolic disease, such as diabetes or obesity,
 CC in a subject. The variation may occur in exons 3, 4 or 10. In exon
 CC 3 variation may be a substitution of a G for a C at nucleotide 320,
 CC resulting in the amino acid substitution P71A; in exon 4 variation may
 CC be a substitution of a T for a C at nucleotide 550; and in exon 10
 CC variation may be a substitution of a T for a C at nucleotide 1037,
 CC resulting in the amino acid substitution R340W. There may also be
 CC nucleotide variation in intron 6. The numbering of these
 CC variations is based on the full length cDNA as given, rather than on
 CC position 1 of the open reading frame.
 XX
 SQ Sequence 1647 BP: 346 A: 502 C: 462 G: 337 T: 0 other;

Query Match 100.0%; Score 101; DB 22; Length 1647;
 Best Local Similarity 100.0%; Pred. No. 1e-22;
 Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gctcaagttcctgcacatcttggtccctgcgcgcgcgcctctctctacgcac 60


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Db 1000 gctcaagttcctgcacatcttggltccctgcgcgcgcgcctcctctctaccgcac 1059
      |||
Oy 61 tatcaagattggcgcacatcgcagacttgctg 101
      |||
Db 1060 tatcaagattggcgcacatcgcagacttgctg 1100
      |||

HRSUT 3
AAD03296
ID AAD03296 standard; DNA: 2109 BP.
AC AAD03296;
XX
XX
XX 13-JUN-2001 (first entry)
XX
XX Human AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.
XX
XX Human: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
KW cystathione beta synthase; CBS; cardiact; gene therapy; ss.
XX Homo sapiens.
OS
XX
XX Key Location/Qualifiers
XX 5'UTR 1..471
XX /tag= a
XX CDS 472..1389
XX /tag= b
XX /product= "Human Prkag3 protein"
XX 3'UTR 1390..2109
XX /tag= c

W0200120003-A2.
XX
XX 22-MAR-2001.
XX
XX 11-SEP-2000; 2000MO-EP09896.
XX
XX 10-SEP-1999; 99EP-0402236.
XX 18-MAY-2000; 2000EP-0401386.
XX
XX (INRG ) INRA INST NAT RECH AGRONOMIQUE.
XX (ANDE/) ANDERSSON L.
XX (LOOF/) LOOFT C.
XX (KALM/) KALM E.
XX
XX Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
XX Iannucciell N, Gellin J, Le Roy P, Chardon P;
XX WPI: 2001-244810/25.
XX P-PSDB: AAE00221.
XX
XX New variants of the gamma subunit of vertebrate adenosine
XX monophosphate-activated kinase for diagnosis or treatment of disorders
XX associated with energy metabolism such as diabetes, obesity, and
XX myopathy.
XX
XX Claim 12; Fig 2; 71pp; English.
XX
XX The present sequence is a cDNA encoding human adenosine monophosphate
XX (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
XX PRKAG3. Mutation in Prkag3 results in an altered regulation of
XX carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
XX useful as therapeutic for treating carbohydrate metabolism disorders such
XX as diabetes, obesity, and disorders associated with muscle metabolism
XX such as myopathy and cardiovascular diseases, to modulate AMPK
XX activity, and for restoring a normal AMPK function. PRKAG3 sequence
XX and its functionally altered mutants are useful for the diagnostic
XX evaluation, genetic testing and prognosis of a metabolic disorder,
XX preferably a carbohydrate metabolism disorder. Primers that can detect
XX a genetic polymorphic marker linked to a sequence encoding PRKAG3, are

```

```

CC useful for detecting a dysfunction of carbohydrate metabolism resulting
CC from the expression of a functionally altered allele of PRKAG3.
CC Transgenic animal and host cell transformed with PRKAG3 or a
CC heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
CC screening compounds able to modulate AMPK activity. Nucleic acid
CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
CC in a sequence encoding the first cystathione beta synthase (CBS) domain
CC of PRKAG3 and is useful in gene therapy.
XX
XX Sequence 2109 BP; 458 A; 621 C; 560 G; 470 T; 0 other;
SQ

Query Match 100.0%; Score 101; DB 22; Length 2109;
Best Local Similarity 100.0%; Pred. No. 1; le-22;
Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 gctcaagttcctgcacatcttggltccctgcgcgcgcgcctcctctaccgcac 60
      |||
Db 900 gctcaagttcctgcacatcttggltccctgcgcgcgcgcctcctctaccgcac 959
      |||
Oy 61 tatcaagattggcgcacatcgcagacttgctg 101
      |||
Db 960 tatcaagattggcgcacatcgcagacttgctg 1000
      |||

RESULT 4
AAD03320
ID AAD03320 standard; cDNA; 2115 BP.
XX
XX AAD03320;
XX
XX 13-JUN-2001 (first entry)
XX
XX Human AMPK gamma subunit muscle-specific isoform, complete PRKAG3 cDNA.
XX
XX Human: gamma subunit; adenosine monophosphate-activated kinase; AMPK;
KW PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
KW genetic testing; carbohydrate metabolism disorder; skeletal muscle;
KW cystathione beta synthase; CBS; cardiact; gene therapy; ss.
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 1..1395
XX /tag= a
XX /product= "Human complete Prkag3 protein"

W0200120003-A2.
XX
XX 22-MAR-2001.
XX
XX 11-SEP-2000; 2000MO-EP09896.
XX
XX 10-SEP-1999; 99EP-0402236.
XX 18-MAY-2000; 2000EP-0401386.
XX
XX (INRG ) INRA INST NAT RECH AGRONOMIQUE.
XX (ANDE/) ANDERSSON L.
XX (LOOF/) LOOFT C.
XX (KALM/) KALM E.
XX
XX Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
XX Iannucciell N, Gellin J, Le Roy P, Chardon P;
XX WPI: 2001-244810/25.
XX P-PSDB: AAE00223.
XX
XX New variants of the gamma subunit of vertebrate adenosine
XX monophosphate-activated kinase for diagnosis or treatment of disorders
XX associated with energy metabolism such as diabetes, obesity, and
XX myopathy.
XX
XX Claim 12; Page 65-68; 71pp; English.
XX

```

XX The present sequence is a cDNA encoding human adenosine monophosphate
 CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
 CC complete PRKAG3. Mutation in Prkag3 results in an altered regulation of
 CC carbohydrate metabolism, particularly in skeletal muscle. PRKAG3 is
 CC useful as therapeutic for treating carbohydrate metabolism disorders such
 CC as diabetes, obesity, and disorders associated with muscle metabolism
 CC such as myopathy and cardiovascular diseases, to modulate AMPK
 CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
 CC and its functionally altered mutants are useful for the diagnostic
 CC evaluation, genetic testing and prognosis of a metabolic disorder,
 CC preferably a carbohydrate metabolism disorder. Primers that can detect
 CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
 CC useful for detecting a dysfunction of carbohydrate metabolism resulting
 CC from the expression of a functionally altered allele of PRKAG3.
 CC Transgenic animal and host cell transformed with PRKAG3 or a
 CC heterotimeric AMPK consisting of PRKAG3 or its mutant, are useful for
 CC screening compounds able to modulate AMPK activity. Nucleic acid
 CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
 CC in a sequence encoding the first cystathione beta synthase (CBS) domain
 CC of PRKAG3 and is useful in gene therapy.

SO Sequence 2115 BP; 460 A; 622 C; 562 G; 471 T; 0 other;

Query Match 100.0%; Score 101; DB 22; Length 2115;
 Best Local Similarity 100.0%; Pred. No. 1.1e-22;
 Matches 101; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 gctcaagttcctgcacatcttggctcctgtgcccggccctcctcctaccgcac 60
 Db 906 gctcaagttcctgcacatcttggctcctgtgcccggccctcctcctaccgcac 965

OY 61 tatccaagattgggcacatccgacattcgagacttgctg 101
 Db 966 tatccaagattgggcacatccgacattcgagacttgctg 1006

RESUI.T 5

AAD03295
 ID AAD03295 standard; cDNA; 1867 BP.

AC AAD03295;

DT 13-JUN-2001 (first entry)

DE Pig AMPK gamma subunit muscle-specific isoform, PRKAG3 cDNA.

KM Pig; gamma subunit; adenosine monophosphate-activated kinase; AMPK;
 KM PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
 KM genetic testing; carbohydrate metabolism disorder; skeletal muscle;
 KM cystathione beta synthase; CBS; cardiant; gene therapy; RN locus;
 KM chromosome 15; ss.

XX Sus scrofa.

OS Key Location/Qualifiers
 FH 1..471
 FT /*tag= a
 FT 472..1389
 FT /*tag= b
 FT /product= "Sus scrofa PRKAG3 protein"
 FT 1390..1867
 FT /*tag= c

PN MO200120003-A2.

PD 22-MAR-2001.

PF 11-SEP-2000; 2000MO-EP09896.

PR 10-SEP-1999; 99EP-0402236.
 PR 18-MAY-2000; 2000EP-0401388.

XX (INRG) INRA INST NAT RECH AGRONOMIQUE.
 PA (ANDE/) ANDERSSON L.
 PA (LOOF/) LOOFT C.
 PA (KALM/) KALM E.
 XX Andersson L, Loof C, Kalm E, Milan D, Robic A, Rogel-Galliard C;
 PI Iannuccelli N, Gellin J, Le Roy P, Chardon P;
 DR WPI; 2001-244810/25.
 XX P-PSDB: AAE00220.

PS Claim 12; Fig 2; 71pp; English.

CC The present sequence is a cDNA encoding pig adenosine monophosphate
 CC (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
 CC PRKAG3. Prkag3 gene is located in the RN locus of chromosome 15.
 CC Mutation in Prkag3 results in an altered regulation of carbohydrate
 CC metabolism, particularly in skeletal muscle. PRKAG3 is useful as
 CC therapeutic for treating carbohydrate metabolism disorders such as
 CC diabetes, obesity, and disorders associated with muscle metabolism
 CC such as myopathy and cardiovascular diseases, to modulate AMPK
 CC activity, and for restoring a normal AMPK function. PRKAG3 sequence
 CC and its functionally altered mutants are useful for the diagnostic
 CC evaluation, genetic testing and prognosis of a metabolic disorder,
 CC preferably a carbohydrate metabolism disorder. Primers that can detect
 CC a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
 CC useful for detecting a dysfunction of carbohydrate metabolism resulting
 CC from the expression of a functionally altered allele of PRKAG3.
 CC Transgenic animal and host cell transformed with PRKAG3 or a
 CC heterotimeric AMPK consisting of PRKAG3 or its mutant, are useful for
 CC screening compounds able to modulate AMPK activity. Nucleic acid
 CC encoding PRKAG3 is useful for detecting mutations in a Prkag3 gene, or
 CC in a sequence encoding the first cystathione beta synthase (CBS) domain
 CC of PRKAG3 and is useful in gene therapy.

SO Sequence 1867 BP; 380 A; 583 C; 529 G; 375 T; 0 other;

Query Match 92.7%; Score 93.6; DB 22; Length 1867;
 Best Local Similarity 96.0%; Pred. No. 2.4e-20;
 Matches 96; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 2 ctcaagttcctgcacatcttggctcctgtgcccggccctcctcctaccgcac 61
 Db 901 ctcaagttcctgcacatcttgggcacctgtgcccggccctcctcctaccgcac 960

OY 62 atccaagattgggcacatccgacattcgagacttgctg 101
 Db 961 atccaagattgggcacatccgacattcgagacttgctg 1000

RESULT 6

AAD03319
 ID AAD03319 standard; cDNA; 1873 BP.

AC AAD03319;

DT 13-JUN-2001 (first entry)

DE Pig AMPK gamma subunit muscle-specific isoform, complete PRKAG3 cDNA.

KM Pig; gamma subunit; adenosine monophosphate-activated kinase; AMPK;
 KM PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
 KM genetic testing; carbohydrate metabolism disorder; skeletal muscle;
 KM cystathione beta synthase; CBS; cardiant; gene therapy; RN locus;
 KM chromosome 15; ss.

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OS   Sus scrofa.
XX   Key      Location/Qualifiers
FH   CDS      1..1395
FT     /*tag= a
FT     /product= "Sus scrofa complete prkag3 protein"
XX   MO200120003-A2.
XX   22-MAR-2001.
XX   11-SEP-2000: 2000WO-EP09896.
XX   10-SEP-1999: 99EP-0402236.
XX   18-MAY-2000: 2000EP-0401388.
XX   (INRG ) INRA INST NAT RECH AGRONOMIQUE.
XX   (ANDE/) ANDERSSON L.
XX   (LOOF/) LOOFT C.
XX   (KALM/) KALM E.
XX   Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
XX   Iannuccelli N, Gellin J, Le Roy P, Chardon P;
XX   WP1: 2001-244810/25.
XX   P-PSDB: AAE00222.
XX   New variants of the gamma subunit of vertebrate adenosine
XX   monophosphate-activated kinase for diagnosis or treatment of disorders
XX   associated with energy metabolism such as diabetes, obesity, and
XX   myopathy -
XX   Claim 12: Page 62-64; 71pp; English.
XX   The present sequence is a cDNA encoding pig adenosine monophosphate
XX   (AMP)-activated kinase (AMPK) gamma subunit muscle-specific isoform,
XX   complete PRKAG3. Prkag3 gene is located in the RN locus of chromosome
XX   15. Mutation in prkag3 results in an altered regulation of carbohydrate
XX   metabolism, particularly in skeletal muscle. PRKAG3 is useful as
XX   therapeutic for treating carbohydrate metabolism disorders such as
XX   diabetes, obesity, and disorders associated with muscle metabolism
XX   such as myopathy and cardiovascular diseases, to modulate AMPK
XX   activity, and for restoring a normal AMPK function. PRKAG3 sequence
XX   and its functionally altered mutants are useful for the diagnostic
XX   evaluation, genetic testing and prognosis of a metabolic disorder.
XX   Preferably a carbohydrate metabolism disorder. Primers that can detect
XX   a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
XX   useful for detecting a dysfunction of carbohydrate metabolism resulting
XX   from the expression of a functionally altered allele of PRKAG3.
XX   Transgenic animal and host cell transformed with PRKAG3 or a
XX   heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
XX   screening compounds able to modulate AMPK activity. Nucleic acid
XX   encoding PRKAG3 is useful for detecting mutations in a prkag3 gene, or
XX   in a sequence encoding the first cystathione beta synthase (CBS) domain
XX   of PRKAG3 and is useful in gene therapy.
XX   Sequence 1873 BP; 382 A; 580 C; 535 G; 376 T; 0 other;
SQ

```

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Query Match      92.7%; Score 93.6; DB 22; Length 1873;
Best Local Similarity 96.0%; Pred. No. 2,4e-20;
Matches 96: Conservative 0; Mismatches 4; Indels 0; Gaps 0;

```

```

OY   2 ctaaagttcctgcacatcttggctccctgctgccccggcccccttccctctacgcagc 61
OY   |||||||
DB   907 ctcaagttcctgcacatcttggcaccctgcgcgcccccttccctctacgcagc 966
OY   |||||||
OY   62 atccaagattggcagtcgacatccgagattgctg 101
OY   |||||||
DB   967 atccaagattggcagtcgacatccgagattgctg 1006

```

RESULT 7

```

AAD03321
ID   AAD03321 standard; DNA: 2022 BP.
XX
XX   AAD03321:
AC   13-JUN-2001 (first entry)
XX
XX   Sus scrofa PRKAG3 splice variant DNA.
XX
XX   Pig: gamma subunit, adenosine monophosphate-activated kinase; AMPK;
XX   PRKAG3; diabetes; obesity; myopathy; cardiovascular disease; anorectic;
XX   genetic testing; carbohydrate metabolism disorder; skeletal muscle;
XX   cystathione beta synthase; CBS; cardiact; gene therapy; ds.
XX   Sus scrofa.
XX
XX   Key      Location/Qualifiers
FH   CDS      1..1345
FT     /*tag= a
FT     /product= "Sus scrofa prkag3 splice variant"
XX   MO200120003-A2.
XX   22-MAR-2001.
XX   11-SEP-2000: 2000WO-EP09896.
XX   10-SEP-1999: 99EP-0402236.
XX   18-MAY-2000: 2000EP-0401388.
XX   (INRG ) INRA INST NAT RECH AGRONOMIQUE.
XX   (ANDE/) ANDERSSON L.
XX   (LOOF/) LOOFT C.
XX   (KALM/) KALM E.
XX   Andersson L, Looft C, Kalm E, Milan D, Robic A, Rogel-Gaillard C;
XX   Iannuccelli N, Gellin J, Le Roy P, Chardon P;
XX   WP1: 2001-244810/25.
XX   P-PSDB: AAE00222.
XX   New variants of the gamma subunit of vertebrate adenosine
XX   monophosphate-activated kinase for diagnosis or treatment of disorders
XX   associated with energy metabolism such as diabetes, obesity, and
XX   myopathy -
XX   Claim 12: Page 69; 71pp; English.
XX   The present sequence is pig adenosine monophosphate (AMP)-activated
XX   kinase (AMPK) gamma subunit muscle-specific isoform, PRKAG3 splice
XX   variant DNA. Prkag3 gene is located in the RN locus of chromosome 15.
XX   Mutation in prkag3 results in an altered regulation of carbohydrate
XX   metabolism, particularly in skeletal muscle. PRKAG3 is useful as
XX   therapeutic for treating carbohydrate metabolism disorders such as
XX   diabetes, obesity, and disorders associated with muscle metabolism
XX   such as myopathy and cardiovascular diseases, to modulate AMPK
XX   activity, and for restoring a normal AMPK function. PRKAG3 sequence
XX   and its functionally altered mutants are useful for the diagnostic
XX   evaluation, genetic testing and prognosis of a metabolic disorder,
XX   preferably a carbohydrate metabolism disorder. Primers that can detect
XX   a genetic polymorphic marker linked to a sequence encoding PRKAG3, are
XX   useful for detecting a dysfunction of carbohydrate metabolism resulting
XX   from the expression of a functionally altered allele of PRKAG3.
XX   Transgenic animal and host cell transformed with PRKAG3 or a
XX   heterotrimeric AMPK consisting of PRKAG3 or its mutant, are useful for
XX   screening compounds able to modulate AMPK activity. Nucleic acid
XX   encoding PRKAG3 is useful for detecting mutations in a prkag3 gene, or
XX   in a sequence encoding the first cystathione beta synthase (CBS) domain
XX   of PRKAG3 and is useful in gene therapy.
XX   Sequence 2022 BP; 412 A; 623 C; 593 G; 394 T; 0 other;
SQ

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Query Match	92.7%	Score 93.6	DB 22	Length 2022
Best Local Similarity	96.0%	Pred. No. 2.4e-20		
Matches 96	Conservative 0	Mismatches 4	Indels 0	Gaps 0
OY	2	ctcaagttccgcgaatcttggttccctgctgcgccgagccctctcctctacgcgact	61	
Db	1057	ctcaagttccgcgaatcttggcacccctgctgcgccgagccctctcctctacgcgacc	116	
OY	62	atccaagattcggagcatcgccacattccgaagactgctg	101	
Db	1117	atccaagattcggagcatcgccacattccgaagactgctg	1156	

RESULT	8
AAH43683	
ID	AAH43683 standard; DNA; 1722 BP.

DT 21-JAN-2002 (first entry)

DE PRKAG3 Intron 4 - Intron 10.

Human; AMP-activated protein kinase gamma 3 subunit; PRKAG3; variant;

OS Homo sapiens.

FH	Key	Location/Qualifiers
EH		

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status = A

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/number= "Exon 5"

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FT      / *tag= c

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FT exon 553...611

Feature	Start	End	Strand	Score	Gene	Transcript	Exon	Intron
Exon 6	612	736	+	1000	FT	FT	612	736

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/number= "Intron
FT

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      / *tag= f
PTT

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FT	Intron	783..986

FT	EST	over	/number="Intron
007	1041		

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/number= "Exon 8"

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1042...1242
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FT exon 1243..1369

/number="Exon 9"

/*tag = k

ET	1523: .1688
exon	/*tag= 1

FT	Exon 10
Intron	1689..1722

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FT
/number= "Intron

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PN W0200177305-A2.

PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-SE00765.
XX
PR 07-APR-2000; 2000US-195665P.

PA (AREX-) AREXIS AB.

PI Andersson L, Luthman H, Marklund S,

DR WPI; 2001-657170/75.

PT New variants of human AMP-activated protein kinase gamma3 subunit
PT associated with a metabolic disease e.g. diabetes or obesity and method
PT for determining a risk estimate of diseases in subject by detecting the
PT variant -

PS Example 1; Fig 3; 25pp; English.

CC The sequences given in AAH343681-84 represents genomic fragments
CC encoding the human AMP-activated protein kinase gamma 3 subunit
CC (PRKAG3). Detecting the presence of the PRKAG3 DNA, or a variant,
CC is useful in determining a risk estimate of a metabolic disease,
CC such as diabetes or obesity, in a subject. The variation may occur
CC in exons 3, 4 or 10. In exon 3 variation may be a substitution of
CC a G for a C at nucleotide 330, resulting in the amino acid
CC substitution P71A. In exon 4 variation may be a substitution of a
CC T for a C at nucleotide 550, and in exon 10 variation may be a
CC substitution of a T for a C at nucleotide 1037, resulting in the
CC amino acid substitution R340W. There may also be nucleotide variation
CC in intron 6.

SQ Sequence 1722 BP; 321 A; 504 C; 534 G; 363 T; 0 other;

Query Match	78.48;	Score	79.2;	DB	22;	Length	1722;
Best Local Similarly	91.38;	Pred. No.	9e-16;				
Matches	84;	Conservative	0;	Mismatches	8;	Indels	0;
				Gaps			0;

Dy 10 ccgcacatcttggttcctcgtgcccggccctctctcctaagacattccaaga 69
 || | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 1510 ccaactaaccaagggttccctcgtgcccggccctctctcctaagacattccaaga 1568

QY	70	tttggcatcggcacatctcggagacttgctg	101
Db	1570	tttggcatcggcacatctcggagacttgctg	1601

RESULT	9
AAx06882	
ID	AAx06882 standard; cDNA; 1435 BP.

AC AAX06882;

DT 26-APR-1999 (first entry)

Disease associated protein kinase DAPK-7 cDNA.

KM	DAMP-7: disease associated protein kinase; human; diagnosis;
KM	thyresap: adult respiratory distress syndrome; allergy; asthma;
KM	arteriosclerosis: bronchitis; emphysema; hyperostonophyllia;
KM	myocardial inflammation; pericardial inflammation; anaemia;
KM	rheumatoid arthritis; Addison's disease; AIDS; atherosclerosis;
KM	atopic dermatitis; dermatomyositis; diabetes mellitus;
KM	glopioculonephritis; gout; Grave's disease; lupus erythematosus;
KM	multiple sclerosis; myasthenia gravis; osteoarthritis;
KM	osteoporosis; pancreatitis; polycystic kidney disease;
KM	polymyositis; scleroderma; Sjogren's syndrome;
KM	autoimmune thyroiditis; cancer; infection; trauma;
KM	cell proliferation; ss.

OS Homo sapiens.

XX 0

FH	Key	Location/Qualifiers
F7	CDS	265..1503
F7P		/*tag= a
XX		
PN	MO9856052-A2.	
XX		
PD	23-DEC-1998.	
XX		
PF	19-JUN-1998;	98MO-US12813.
XX		
PB	19-JUN-1997;	97US-0878989.
XX		
PA	(INCY-) INCYTE PHARM INC.	
PI	Bandman O., Corley NC, Goli SK, Guegler KJ, Hillman JL;	
PI	Lai P., Shah P;	
DR	WPI: 1999-080952/07.	
DR	P-PSDB: AAW88438.	
XX		
PT	New disease associated protein kinases - used to stimulate cell	
PT	proliferation and to treat the immune response and cancer	
XX		
XS	Claim 5; Page 66-67; 93pp; English.	

This cDNA sequence codes for human disease associated protein kinase DAPK7 (see AAM68438). DAPK-7 cDNA was first identified in the PENTUT01 cDNA library using a computer search for amino acid alignments and a consensus sequence was derived from the extended and overlapping Inocyte clones 3075712/HENR0T01, 842220/PROSTU05, 1364747/SCORNO02, 1459572 and 145802/PENTUT01 and 1479332/CORNOT02. DAPK-7 shows 73% homology with the human foetal liver AMPK gamma subunit (GI 1335656), and is associated with cDNA libraries which are immortalised or cancerous and show inflammatory or immune responses. The invention provides disease associated protein kinases DAPK-1 to DAPK-7 (see AAM68432-38) and cDNA clones encoding them (see AAX06831-36 and AAX06882), as well as expression vectors, host cells, agonists, antagonists and antibodies. The invention further provides uses of such products in the diagnosis, prevention and treatment of diseases associated with cell proliferation, especially cancer or an immune response (claimed). Conditions that may be treated include adult respiratory distress syndrome, allergies, asthma, arteriosclerosis, bronchitis, emphysema, hyperostinophilia, myocardial or pericardial inflammation, rheumatoid arthritis, Addison's disease, AIDS, anaemia, atherosclerosis, various diseases of the digestive system, atopic dermatitis, dermatomyositis, diabetes mellitus, glomerulonephritis, gout, Grave's disease, lupus erythematosus, multiple sclerosis, myasthenia gravis, osteoarthritis, osteoporosis, pancreatitis, polycystic kidney disease, polymyositis, scleroderma, Sjogren's syndrome, autoimmune thyroiditis, complications of cancer, extracorporeal circulation, viral, bacterial, fungal, parasitic, protozoal and helminthic infections, and trauma (disclosed). The DAPK nucleic acids are also used in a method for detection of DAPK expression levels in a biological sample.

Sequence 1435 BP; 421 A; 298 C; 331 G; 385 T; 0 other;

Query Match	30.5%;	Score 30.8;	DB 20;	Length 1435;
Best Local Similarity	57.1%;	Pred. No. 2.3;		
Matches 56;	Conservative	0;	Mismatches 42;	Indels 0;
				Gaps 0

2 ctcaagatctccgcagacatcttggttcaccgcgtcccgagccctacttctctaccgaact 61
 ||||| ||||| || ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 617 ctcaagatctccgcagacatcttggttcaccgcgtcccgagccctacttctctaccgaac 676
 62 atccaagaatttggcgcacgcgcacatctccgaagacttgcg 99
 || ||| || ||| || ||| || ||| || ||| || ||| || ||| || ||| || ||| || |||
 677 ctgaatcagctctggaatcagaacgctaccacaacatctgc 714

RESULT 10

AA584265	AA584265 standard; cDNA; 1467 BP.
AC	AA584265;
DT	13-FEB-2002 (first entry)
DE	DNA encoding novel human diagnostic protein #20069.
KW	Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX	food supplement; medical imaging; diagnostic; genetic disorder; ss.
OS	Homo sapiens.
PN	MO200175067-A2.
PD	11-OCT-2001.
PF	30-MAR-2001; 2001WO-US08631.
PR	31-MAR-2000; 2000US-0540217.
XX	23-AUG-2000; 2000US-0649167.
PA	(HYSE-) HYSEQ INC.
PI	Drmanac RT, Liu C, Tang YT;
DR	WPI: 2001-639362/73.
XX	P-PSDB; ABG20078.
PT	New isolated polynucleotide and encoded polypeptides, useful in
PT	diagnostics, forensics, gene mapping, identification of mutations
PT	responsible for genetic disorders or other traits and to assess
PT	biodiversity -

PS Claim 1 SEQ ID No 20069: 103pp:English.

xx

CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. MS64197-AB594564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at [ftp.wipo.int/publ/published_pcl_sequences](http://wipo.int/publ/published_pcl_sequences).

SQ Sequence 1467 BP; 416 A; 327 C; 343 G; 381 T; 0 other;

Query Match	30.5%	Score 30.8;	DB 23;	Length 1467;
Best Local Similarity	57.1%;	Pred. No. 2.3;		
Matches 56;	Conservative	0;	Mismatches 42;	Indels 0;
				Gaps 0

[illegible]

RESULT 11

AAH14839 ID AAH14839 standard; cDNA: 2223 BP.

XX AC AAH14839;

XX DT 26-JUN-2001 (first entry)

XX XX

DE Human cDNA sequence SEQ ID NO:12660.

OS Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss

XX Homo sapiens.

PN EP1074617-A2.

PD 07-FEB-2001.

XX 28-JUL-2000; 2000EP-0116126.

XX 29-JUL-1999; 99JP-0248036.

PR 27-AUG-1999; 99JP-0300253.

PR 11-JAN-2000; 2000JP-0118776.

PR 02-MAY-2000; 2000JP-0183767.

PR 09-JUN-2000; 2000JP-0241899.

PA (HELI-) HELIX RES INST.

PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

DR WPI: 2001-318749/34.

XX

PT Primer sets for synthesizing polynucleotides, particularly the 5602

PT full-length cDNAs defined in the specification, and for the detection

PT and/or diagnosis of the abnormality of the proteins encoded by the

PT full-length cDNAs -

XX

XX Claim 8; SEQ ID 12660; 2537bp + CD ROM; English.

PS

XX The present invention describes primer sets for synthesizing 5602

CC full-length cDNAs defined in the specification. Where a primer set

CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary

CC to the complementary strand of a polynucleotide which comprises one of

CC the 5602 nucleotide sequences defined in the specification, where the

CC of an oligonucleotide comprising at least 15 nucleotides; or (b) a combination

CC complementary strand of a polynucleotide which comprises a 5'-end

CC sequence and an oligonucleotide comprising a sequence complementary to a

CC polynucleotide which comprises a 3'-end sequence, where the

CC the 5'-end sequence/3'-end sequence is selected from those defined in

CC the specification. The primer sets can be used in antisense therapy and

CC in gene therapy. The primers are useful for synthesizing polynucleotides,

CC particularly full-length cDNAs. The primers are also useful for the

CC detection and/or diagnosis of the abnormality of the proteins encoded by

CC the full-length cDNAs. The primers allow obtaining of the full-length

CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and

CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to

CC AAB93893 represent human amino acid sequences; and AAH13629 to AAH13632

CC represent oligonucleotides, all of which are used in the exemplification

XX of the present invention.

XX

XX Sequence 2223 BP: 633 A: 431 C: 496 G: 663 T: 0 other;

XX

Query Match 30.5% Score 30.8; DB 22; Length 2223;

Best Local Similarity 57.1%; Pred. No. 2.5;

Matches 56; Conservative 0; Mismatches 42; Indels 0; Gaps 0;

2 ctcaagttctcgacactcttgctccctgctgcgccgcgcctctctctcaacgcgact 61

Dy
62 atccaaatttgcgcatcgacatcgcgaactggc 99

Dd
709 ctfgatgagcttgtaataagaacgtaccacaacatlgc 746

RESULT	12
ID	AAS05559/c
standard:	DNA; 566 BP.
XX	
AC	AAS05559;
XX	
DT	07-SEP-2001 (first entry)
XX	
DE	Mammalian vestibular system geotactic behaviour modulator gene #159.
XX	
KM	Mammalian vestibular system: invertebrate; geotactic behaviour: vertigo;
KW	graviperceptive disorder; motion sickness; labyrinthitis; syphilis; de;
KM	Meniere's disease; acoustic neuroma; multiple sclerosis; epilepsy;
KM	trauma; infection of the middle ear; ototoxic agent exposure.
XX	
OS	Drosophila melanogaster .
XX	
PN	MO200140519-A2.
XX	
PD	07-JUN-2001.
XX	
PF	01-DEC-2000; 2000WO-US32639.
XX	
PR	02-DEC-1999; 99US-0168579.
XX	
PA	26-SEP-2000; 2000US-0669751.
XX	
PI	(NEUR-) NEUROSCIENCES RES FOUND INC.
XX	
PI	Greenspan RJ;
XX	
DR	WPI; 2001-356159/37.
XX	
PT	New isolated nucleic acid having mammalian vestibular system-modulating
PT	activity useful in the treatment of disorders such as motion sickness
PT	and vertigo -
XX	
PS	Claim 59; Page 144; 179pp; English.
XX	
CC	The sequences shown in AAS05401-AAS05661 represent DNA with mammalian
CC	vestibular system-modulating activity. The DNA sequences can be used in a
CC	method whereby a first and second strain of an invertebrate is obtained,
CC	and both are subjected to conditions in which the strains exhibit
CC	different geotactic behaviour. Genes that are differentially expressed in
CC	the first strain relative to the second strain are then identified.
CC	Mammalian genes having substantially the same nucleic acid sequence as
CC	these genes are used to decrease the symptoms of graviperceptive
CC	disorders such as motion sickness, vertigo, labyrinthitis, Meniere's
CC	disease, acoustic neuroma, multiple sclerosis, syphilis, trauma,
CC	infection of the middle ear, exposure to ototoxic agents and epilepsy.
XX	
SQ	Sequence 566 BP; 191 A; 129 C; 161 G; 85 T; 0 other;

Query Match	29.3%; Score 29.6; DB 22; Length 566;
Best Local Similarity	68.3%; Pred No. 4.4;
Matches 41; Conservative 0; Mismatches 19; Indels 0; Gaps 0;	

OY	13 gcaacatcttggtcctgcgcccgcgccttccctaccgaactlccaagattt 72
Db	544 GCCTATCAAGGCGACTACTGTCTGCCTCTTCTTCTTAACCTCATTTCCCTGATC 485

RESULT	13
AAAC37238	

ID AAC37238 standard: DNA: 1326 BP.
XX AAC37238;
AC
XX 17-OCT-2000 (first entry)
DT
XX
DE Arabidopsis thaliana DNA fragment SEQ ID NO: 16667.
XX
XX Hybridisation assay; genetic mapping; gene expression control;
KM protein identification; signal transduction pathway;
KW metabolic pathway; promoter; termination sequence; ss.
XX
OS Arabidopsis thaliana.
PM EP1033405-A2.
XX
PD 06-SEP-2000.
XX
PF 25-FEB-2000; 2000EP-0301439.
XX
PR 25-FEB-1999; 99US-0121825.
PR 05-MAR-1999; 99US-0123180.
PR 09-MAR-1999; 99US-0123548.
PR 23-MAR-1999; 99US-0125788.
PR 25-MAR-1999; 99US-0126264.
PR 29-MAR-1999; 99US-0126785.
PR 01-APR-1999; 99US-0127462.
PR 06-APR-1999; 99US-0128234.
PR 08-APR-1999; 99US-0128714.
PR 16-APR-1999; 99US-0129845.
PR 19-APR-1999; 99US-0130077.
PR 21-APR-1999; 99US-0130449.
PR 23-APR-1999; 99US-0130510.
PR 28-APR-1999; 99US-0130891.
PR 30-APR-1999; 99US-0131449.
PR 30-APR-1999; 99US-0132048.
PR 04-MAY-1999; 99US-0132407.
PR 05-MAY-1999; 99US-0132484.
PR 06-MAY-1999; 99US-0132485.
PR 06-MAY-1999; 99US-0132486.
PR 07-MAY-1999; 99US-0132487.
PR 11-MAY-1999; 99US-0132863.
PR 14-MAY-1999; 99US-0134256.
PR 14-MAY-1999; 99US-0134218.
PR 14-MAY-1999; 99US-0134219.
PR 14-MAY-1999; 99US-0134221.
PR 18-MAY-1999; 99US-0134370.
PR 19-MAY-1999; 99US-0134768.
PR 20-MAY-1999; 99US-0135124.
PR 21-MAY-1999; 99US-0135124.
PR 24-MAY-1999; 99US-0135529.
PR 25-MAY-1999; 99US-0135629.
PR 27-MAY-1999; 99US-0136021.
PR 28-MAY-1999; 99US-0136392.
PR 01-JUN-1999; 99US-0136782.
PR 03-JUN-1999; 99US-0137222.
PR 04-JUN-1999; 99US-0137528.
PR 07-JUN-1999; 99US-0137502.
PR 08-JUN-1999; 99US-0137724.
PR 10-JUN-1999; 99US-0138094.
PR 10-JUN-1999; 99US-0138540.
PR 14-JUN-1999; 99US-0138847.
PR 16-JUN-1999; 99US-0139119.
PR 16-JUN-1999; 99US-0139452.
PR 17-JUN-1999; 99US-0139453.
PR 18-JUN-1999; 99US-0139492.
PR 18-JUN-1999; 99US-0139494.
PR 18-JUN-1999; 99US-0139456.
PR 18-JUN-1999; 99US-0139457.
PR 18-JUN-1999; 99US-0139458.
PR 18-JUN-1999; 99US-0139459.
PR 18-JUN-1999; 99US-0139460.

PR 18-JUN-1999; 99US-0139461.
PR 18-JUN-1999; 99US-0139462.
PR 18-JUN-1999; 99US-0139463.
PR 18-JUN-1999; 99US-0139750.
PR 18-JUN-1999; 99US-0139763.
PR 21-JUN-1999; 99US-0139817.
PR 22-JUN-1999; 99US-0139899.
PR 23-JUN-1999; 99US-0140353.
PR 23-JUN-1999; 99US-0140354.
PR 24-JUN-1999; 99US-0140354.
PR 28-JUN-1999; 99US-0140695.
PR 29-JUN-1999; 99US-0140823.
PR 30-JUN-1999; 99US-0140991.
PR 01-JUL-1999; 99US-0141287.
PR 01-JUL-1999; 99US-0141842.
PR 02-JUL-1999; 99US-0142154.
PR 06-JUL-1999; 99US-0142055.
PR 08-JUL-1999; 99US-0142390.
PR 09-JUL-1999; 99US-0142803.
PR 12-JUL-1999; 99US-0142920.
PR 13-JUL-1999; 99US-0143542.
PR 14-JUL-1999; 99US-0143624.
PR 15-JUL-1999; 99US-0144005.
PR 16-JUL-1999; 99US-0144085.
PR 16-JUL-1999; 99US-0144086.
PR 19-JUL-1999; 99US-0144325.
PR 19-JUL-1999; 99US-0144331.
PR 19-JUL-1999; 99US-0144332.
PR 19-JUL-1999; 99US-0144333.
PR 19-JUL-1999; 99US-0144334.
PR 19-JUL-1999; 99US-0144335.
PR 20-JUL-1999; 99US-0144352.
PR 20-JUL-1999; 99US-0144632.
PR 21-JUL-1999; 99US-0144684.
PR 21-JUL-1999; 99US-0144814.
PR 21-JUL-1999; 99US-0145086.
PR 22-JUL-1999; 99US-0145088.
PR 22-JUL-1999; 99US-0145087.
PR 22-JUL-1999; 99US-0145089.
PR 22-JUL-1999; 99US-0145192.
PR 23-JUL-1999; 99US-0145145.
PR 23-JUL-1999; 99US-0145218.
PR 23-JUL-1999; 99US-0145224.
PR 26-JUL-1999; 99US-0145276.
PR 27-JUL-1999; 99US-0145913.
PR 27-JUL-1999; 99US-0145918.
PR 28-JUL-1999; 99US-0145919.
PR 28-JUL-1999; 99US-0145951.
PR 02-AUG-1999; 99US-0146386.
PR 02-AUG-1999; 99US-0146388.
PR 03-AUG-1999; 99US-0146389.
PR 03-AUG-1999; 99US-0147038.
PR 04-AUG-1999; 99US-0147204.
PR 05-AUG-1999; 99US-0147302.
PR 05-AUG-1999; 99US-0147192.
PR 06-AUG-1999; 99US-0147260.
PR 06-AUG-1999; 99US-0147303.
PR 09-AUG-1999; 99US-0147416.
PR 09-AUG-1999; 99US-0147493.
PR 09-AUG-1999; 99US-0147935.
PR 10-AUG-1999; 99US-0148171.
PR 11-AUG-1999; 99US-0148319.
PR 12-AUG-1999; 99US-0148341.
PR 13-AUG-1999; 99US-0148565.
PR 13-AUG-1999; 99US-0148684.
PR 16-AUG-1999; 99US-0149368.
PR 17-AUG-1999; 99US-0149175.
PR 18-AUG-1999; 99US-0149426.
PR 20-AUG-1999; 99US-0149722.
PR 20-AUG-1999; 99US-0149723.
PR 20-AUG-1999; 99US-0149929.
PR 23-AUG-1999; 99US-0149902.


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PD 27-SEP-2001.
XX
XX 23-MAR-2001; 2001WO-US09231.
XX
XX 23-MAR-2000; 2000US-191637P.
XX
XX 11-JUL-2000; 2000US-0614150.
XX
XX (PEKE ) PE CORP NY.
XX
XX Ventler JC, Adams M, Li PWD, Myers EW;
XX
XX WPI; 2001-656860/75.
XX
XX P-PSDB; ABB64095.
XX
XX New isolated nucleic acid detection reagent for detecting 1000 or more
XX genes from Drosophila and for elucidating cell signalling and cell-cell
XX interactions -
XX
XX Claim 1: SEQ ID NO 19076; 21pp + Sequence Listing; English.
XX
XX The invention relates to an isolated nucleic acid detection reagent
XX capable of detecting 1000 or more genes from Drosophila. The invention is
XX useful in developmental biology and in elucidating cell signalling and
XX cell-cell interactions in higher eukaryotes for the development of
XX insecticides, therapeutics and pharmaceutical drugs. The invention
XX discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
XX sequences (ABL101840-ABL16175) and the encoded proteins
XX (ABB57737-ABB72072).
XX
XX The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 4363 BP; 1461 A; 939 C; 974 G; 989 T; 0 other;
XX
XX
XX Query Match 29.3%; Score 29.6; DB 23; Length 4363;
XX Best Local Similarity 68.3%; Pred. No. 7;
XX Matches 41; Conservative 0; Mismatches 19; Indels 0; Gaps 0;
XX
XX 13 gcaatcttggtccctcgtgcgcgcgcgcctcctcctctacgacataccaagattt 72
XX ||||||| ||| | ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
XX Db 1544 GCTCATCAAGGCTACTACTGTTCTGCTCTTCTCTCTACCTCATATCCCTGATCT 1485

```

Search completed: October 3, 2002, 16:30:48
 Job time: 14318 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:15:08 ; Search time 5701.1 Seconds
(without alignments)
239.110 Million cell updates/sec

Title: US-09-826-581-5_COPY_1000_1100
Perfect score: 101
Sequence: 1 gcccaagtcctgcacatc.....cacatccgagacttgctg 101

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estln:*
4: em_estmu:*
5: em_estcov:*
6: em_estcpl:*
7: em_estlro:*
8: em_hlc:*
9: gb_est1:*
10: gb_est2:*
11: gb_hlc:*
12: gb_gss:*
13: em_gss_hum:*
14: em_gss_inv:*
15: em_gss_pln:*
16: em_gss_vrl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	93.6	92.7	572	10	B1344527 373008 MA
2	59	58.4	413	9	AA178898 zp38d10. r
3	48.8	48.3	633	10	B1072114 B1072114
4	38.8	38.4	514	10	BC037921 d66908.Y
5	34.8	34.5	918	10	BC027175 602295858
6	32.2	31.9	451	9	AM147353 da01h1.Y
7	32.2	31.9	470	10	B1036397 B1036397
8	32.2	31.9	580	9	AV914030 AV914030
9	31.4	31.1	684	10	BC101440 y72c12.Y
10	31.2	30.9	536	10	BM488662 psm2n.pk0
11	31.2	30.9	595	10	BM487789
12	31.2	30.9	647	10	BM440762
13	31.2	30.9	649	9	AJ395115
14	31.2	30.9	758	9	AJ396118
15	30.8	30.5	413	10	H64260 yf70c12.r1
16	30.8	30.5	451	10	W39604 zc19b04.r1
17	30.8	30.5	474	10	BE803695

18	30.8	30.5	506	10	B1760420
19	30.8	30.5	525	10	BG148719
20	30.8	30.5	601	10	BE846246
21	30.8	30.5	764	9	AU138307
22	30.8	30.5	849	10	B1657718
23	30.8	30.5	886	10	BE572824
24	30.8	30.5	909	9	AL558594
25	30.8	30.5	951	9	AL513784
26	30.8	30.5	967	9	AL524822
27	30.4	30.1	312	10	C06172
28	30.2	29.9	924	12	CNS029MX
29	30	29.7	663	10	BG415737
30	29.8	29.5	784	9	AU080009
31	29.8	29.5	964	10	BC035134
32	29.6	29.3	322	9	AV525978
33	29.6	29.3	619	10	BE978092
34	29.6	29.3	626	10	BF528264
35	29.6	29.3	715	10	BF488250
36	29.6	29.3	754	9	A1063338
37	29.6	29.3	836	10	B1659310
38	29.4	29.1	388	12	AO535079
39	29.4	29.1	559	9	A1062936
40	29.4	29.1	559	9	A1062937
41	29.4	29.1	588	10	BM340317
42	29.4	29.1	699	12	BM447226
43	29.4	29.1	888	9	AL581895
44	29.2	28.9	224	9	AA045087
45	29.2	28.9	388	10	B1774231

ALIGNMENTS

RESULT 1
LOCUS B1344527 572 bp mRNA linear EST 30-JUL-2001
DEFINITION 373008 MARC 2P1G Sus scrofa cDNA 5', mRNA sequence.
VERSION B1344527
KEYWORDS B1344527.1 G1:15037807
SOURCE EST.
ORGANISM Sus scrofa
PLG
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Sus.

REFERENCE

1 (bases 1 to 572)
Fahrenkrug,S.C., Freking,B.A., Rohrer,G.A., Smith,T.P.L., Casas,E.,
Stone,R.T., Heaton,M.P., Grosse,W.M., Bennett,G.A., Laegreid,W.W.
and Keeler,J.W.
Design and use of two pooled tissue normalized cDNA libraries for
EST discovery in swine

JOURNAL

Unpublished (2000)
Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390

Email: smith@mail.marc.usda.gov
Single pass sequencing. Bases called and alt-trimmed with phred
v0.980904.e. Vector identified by cross-match with the -minscore 18
and -mismatch 12 options.

PCR Primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCGCCAGTCACACG
Plate: 119 row: 1 column: 11
Seq primer: ATTGAGTGACACTATAG.
Location/Qualifiers
1..572

FEATURES

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/db_xref="taxon:9823"
/clone_lib="MARC 2P1G"
/tissue_type="pooled"
/lab_host="DH10B"
/note="Vector: pCMV SPORT6; Site_1: XbaI; Site_2: XhoI;

Library made from pooled tissue from testis, ovary, endometrium, hypothalamus, pituitary, and placenta.
BASE COUNT 106 a 190 c 151 g 124 t 1 others
ORIGIN

Query Match 92.7%: Score 93.6; DB 10; Length 572;
Best Local Similarity 96.0%; Pred. No. 7.9e-16;
Matches 96; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
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Db 163 CTCACGTCCTCCGACATCTTGGCACCCCTGCCCCCTCTCTACCGACC 222
CTCACGTCCTCCGACATCTTGGCACCCCTGCCCCCTCTCTACCGACC 222
OY 62 atccaagattggcgcacatccgacatccgacatctgctg 101
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Db 223 ATCCAGATTTGGGCAATCGCACATTCGAGACTTGGCGC 262
ATCCAGATTTGGGCAATCGCACATTCGAGACTTGGCGC 262

RESULT 2
AA178898 413 bp mRNA linear EST 09-MAR-1998
LOCUS 2938010.r1 StrataGene muscle 937209 Homo sapiens cDNA clone
DEFINITION IMAGE:611731 5' similar to SW:AAKG_RAT P80385 5'-AMP-ACTIVATED
PROTEIN KINASE, GAMMA CHAIN /, mRNA sequence.
AA178898
VERSION AA178898.1 GI:1760259
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi;
Mammalia: Eutheria: Primates: Catarrhini: Homiidae: Homo.
AUTHORS 1 (bases 1 to 413)
Hillier, L., Allen, M., Bowles, L., Dubuque, T., Giesel, G., Jost, S.,
Kizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin,
J., Moore, B., Scheinberg, K., Steptoe, M., Tan, F., Theising, B.,
White, Y., Wylie, T., Waterston, R. and Wilson, R.
WashU-NCI human EST Project
TITLE Unpublished (1997)
JOURNAL Contact: Wilson RK
COMMENT Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.wustl.edu
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1280 Std Error: 0.00
Seq primer: -28M13 rev2 from AmerSham
High quality sequence stop: 255.
Location/Qualifiers
1. 413

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/db_xref="taxon:9606"
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/clone_lib="StrataGene muscle 937209"
/tissue_type="muscle"
/dev_stage="adult"
/lab_host="SOLR (kanamycin resistant)"
/note="Organ: skeletal muscle; Vector: pBluescript SK-;
Site_1: EcoRI; Site_2: XhoI; Cloned unidirectionally.
Primer: Oligo dT. Skeletal muscle from patient with
malignant hyperthermia. Average insert size: 1.0 kb;
Uni-ZAP XR Vector; -5' adaptor sequence: 5' GAATTCGCGACAG
3' -3' adaptor sequence: 5' CTCACGTCCTTCTTTTCTTTTCTTTT 3' -

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ORIGIN

Query Match 58.4%; Score 59; DB 9; Length 413;
Best Local Similarity 100.0%; Pred. No. 2.1e-06;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 CTCCTCTCTTACCGACATATCCAGATTTGGGCAATCGCACATTCGAGACTTGGCTG 59
CTCCTCTCTTACCGACATATCCAGATTTGGGCAATCGCACATTCGAGACTTGGCTG 59

RESULT 3
BJ072114 633 bp mRNA linear EST 11-DEC-2001
LOCUS BJ072114 NIBB Mochii normalized Xenopus tailbud library Xenopus
DEFINITION laevis cDNA clone X1096j16 5', mRNA sequence.
BJ072114
ACCESSION BJ072114.1 GI:17502303
VERSION BJ072114.1 GI:17502303
KEYWORDS EST.
SOURCE African clawed frog.
Xenopus laevis
ORGANISM Xenopus laevis

REFERENCE Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi;
Amphibia: Batrachia: Anura: Mesobatrachia: Pipidoidea: Pipidae;
Xenopodinae: Xenopus.
1 (bases 1 to 633)
Kitayama, A., Terasaka, C., Mochii, M., Ueno, N., Shin-I, T. and Kohara,
Y.
Expressed genes in X. laevis embryo
Unpublished (2001)
CONTACT: Tadasu Shin-I
Center for Genetic Resource Information
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tshin@genes.nig.ac.jp.

FEATURES
source
Location/Qualifiers
1. 633
/organism="Xenopus laevis"
/db_xref="taxon:8355"
/clone="X1096j16"
/clone_lib="NIBB Mochii normalized Xenopus tailbud
library"
/tissue_type="whole embryo"
/dev_stage="stage 25"
BASE COUNT 144 a 162 c 148 g 179 t
ORIGIN

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Best Local Similarity 68.0%; Pred. No. 0.0015;
Matches 68; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

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|||||
Db 131 CTCACGTCCTCCGACATCTTGGCACCCCTGCCCCCTCTCTACCGACC 190
CTCACGTCCTCCGACATCTTGGCACCCCTGCCCCCTCTCTACCGACC 190
OY 62 atccaagattggcgcacatccgacatccgacatctgctg 101
|||||
Db 191 ATCTGAGCTGGGCGCATCGCACATTCAGAGATATAGCAG 230
ATCTGAGCTGGGCGCATCGCACATTCAGAGATATAGCAG 230

RESULT 4
BG037921 514 bp mRNA linear EST 24-JAN-2001
LOCUS BG037921 NICHG XGC Emb1 Xenopus laevis cDNA clone IMAGE:3402446
DEFINITION 5' similar to TR:Q99LX8 Q99LX8 H91620P.;, mRNA sequence.
BG037921
ACCESSION BG037921.1 GI:12480506
VERSION BG037921.1 GI:12480506
KEYWORDS EST.
SOURCE African clawed frog.
Xenopus laevis
ORGANISM Xenopus laevis

REFERENCE Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi;
Amphibia: Batrachia: Anura: Mesobatrachia: Pipidoidea: Pipidae;
Xenopodinae: Xenopus.
1 (bases 1 to 514)
NCI-CCAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Query Match	31.9%	Score 32.2;	DB 10;	Length 470;
Best Local Similarity	61.2%;	Pred. No. 47;		
Matches	52;	Mismatches	33;	Indels 0; Gaps 0;
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RESULT	9
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LOCUS	BGI01440
DEFINITION	684 bp mRNA linear EST 29-JAN-2001 uy72c12.y1 McCarrey Eddy round spermatid Mus musculus cDNA clone IMAGE:3664919 5' similar to TR:Q9ULX8 Q9ULX8 H9I620P. ;, mRNA sequence.
ACCESSION	BGI01440
VERSION	BGI01440.1 GI:12596757
KEYWORDS	EST.
SOURCE	house mouse.
ORGANISM	Mus musculus
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus;

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: October 3, 2002, 16:24:51 ; Search time 180.77 Seconds
(without alignments)
137.241 Million cell updates/sec

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Perfect score:

Scoring table: IDENTITY_NUC
Gapop 10.0 ; Gapext 1.0

Searched: 383533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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6: /cgn2_6/ptodata/2/1na/backfile1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	30.8	30.5	1435	2 US-08-878-989-14	Sequence 14, Appl
2	30.8	30.5	1435	4 US-09-272-796-14	Sequence 14, Appl
3	29.4	29.1	1260	1 US-07-866-979-3	Sequence 3, Appl
4	29.4	29.1	1260	2 US-08-466-9068-3	Sequence 3, Appl
5	29.4	29.1	1260	3 US-08-706-281A-3	Sequence 3, Appl
6	29.4	29.1	1260	4 US-09-201-746-3	Sequence 3, Appl
7	29.4	29.1	1260	4 US-09-097-231-3	Sequence 3, Appl
8	27.8	27.5	1378	1 US-08-759-848-2	Sequence 2, Appl
9	27.8	27.5	1378	5 PCT-US95-09383-2	Sequence 2, Appl
10	27.6	27.3	5020	3 US-08-938-291A-3	Sequence 3, Appl
11	26.4	26.1	5975	1 US-08-336-257A-3	Sequence 3, Appl
12	25.6	25.3	1984	2 US-08-822-028-16	Sequence 3, Appl
13	25.6	25.3	1984	4 US-08-479-285-16	Sequence 16, Appl
14	25.4	25.1	1509	2 US-08-481-337A-1	Sequence 1, Appl
15	25.4	25.1	1509	2 US-08-696-268A-1	Sequence 1, Appl
16	25.4	25.1	1509	5 PCT-US95-05467-1	Sequence 1, Appl
17	25.4	25.1	1596	5 PCT-US94-11328A-3	Sequence 1, Appl
18	25.4	25.1	1702	4 US-08-934-254-26	Sequence 26, Appl
19	25.4	25.1	1984	4 US-09-382-256-1	Sequence 1, Appl
20	25.4	25.1	1984	4 US-09-395-115-1	Sequence 1, Appl
21	25.4	25.1	1984	4 US-08-436-265-1	Sequence 1, Appl
22	25.2	25.0	2264	4 US-09-126-109-9	Sequence 9, Appl
23	25.2	25.0	5962	6 5386025-5	Sequence 1, Appl
24	25.2	25.0	5975	1 US-08-404-354B-1	Sequence 1, Appl
25	25.2	25.0	5975	1 US-08-314-083B-1	Sequence 1, Appl
26	25.2	25.0	5975	1 US-08-435-675B-1	Sequence 1, Appl
27	25.2	25.0	5975	3 US-08-884-599-1	Sequence 1, Appl

C 28	24.8	24.6	1419	1 US-08-103-739B-1	Sequence 1, Appl
C 29	24.8	24.6	1419	1 US-08-474-404-1	Sequence 1, Appl
C 30	24.8	24.6	1419	2 US-08-485-845-1	Sequence 1, Appl
C 31	24.8	24.6	1419	2 US-08-482-714-1	Sequence 1, Appl
C 32	24.8	24.6	1419	4 US-09-211-416-1	Sequence 1, Appl
C 33	24.8	24.6	1419	4 US-09-059-958-1	Sequence 1, Appl
C 34	24.4	24.2	261	2 US-08-592-383-7	Sequence 7, Appl
C 35	24.4	24.2	1071	1 US-08-612-986-1	Sequence 1, Appl
C 36	24.4	24.2	1071	1 US-08-361-806A-1	Sequence 1, Appl
C 37	24.4	24.2	1071	5 PCT-US95-16806A-1	Sequence 1, Appl
C 38	24.4	24.2	1094	2 US-08-902-294-1	Sequence 1, Appl
C 39	24.4	24.2	1094	3 US-09-178-637-1	Sequence 1, Appl
C 40	24.4	24.2	2658	2 US-08-592-383-3	Sequence 1, Appl
C 41	24.4	24.2	2928	2 US-08-095-728B-3	Sequence 3, Appl
C 42	24.4	24.2	2928	5 PCT-US92-02320A-3	Sequence 3, Appl
C 43	24.4	24.2	2940	2 US-08-592-383-1	Sequence 3, Appl
C 44	24.4	24.2	2940	6 5171671-1	Sequence 1, Appl
C 45	24.4	24.2	4002	1 US-08-331-488A-1	Sequence 1, Appl

ALIGNMENTS

RESULT 1
US-08-878-989-14
; Sequence 14, Application US/08878989
; Patent No. 5885803
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Corley, Neil C.
; APPLICANT: Guegler, Karl G.
; APPLICANT: Lal, Preeti
; APPLICANT: Goli, Surya K.
; APPLICANT: Shah, Purvi
; TITLE OF INVENTION: DISEASE ASSOCIATED PROTEIN
; NUMBER OF INVENTION: KINASES
; NUMBER OF SEQUENCES: 21
; CORRESPONDENCE ADDRESS:
; ADDRESS: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/878,989
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0321 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; TELEX:
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1435 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: PENITOT01

CLONE: 1452972
US-08-878-989-14

Query Match 30.5%: Score 30.8; DB 2; Length 1435;
Best Local Similarity 57.1%: Pred. No. 0.11;
Matches 56: Conservative 0; Mismatches 42; Indels 0; Gaps 0;

OY 2 ctcaagttcctgacacatcttggtccctgcgcgcgcctctctctacgcact 61
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Db 617 CTCAGTTCCTCCACACTTTTATGTCATATGCCAAGCCTGCTTCATGAGCAGAAC 676
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OY 62 atccaagattggcattcgacatccgagactgac 99
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Db 677 CTGATGAGCTTGGAATGAGAACGTACCAACAACTTGC 714
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RESULT 2
US-09-272-796-14
Sequence 14, Application US/09272796
Patent No. 6207148
GENERAL INFORMATION:
APPLICANT: Bandman, Olga
APPLICANT: Hillman, Jennifer L.
APPLICANT: Corley, Neil C.
APPLICANT: Guegler, Karl G.
APPLICANT: Lal, Surya K.
APPLICANT: Shah, Purvi
TITLE OF INVENTION: DISEASE ASSOCIATED PROTEIN
NUMBER OF SEQUENCES: 21
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/272,796
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/878,989
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0321 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 1435 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: PENITUT01
CLONE: 1452972
US-09-272-796-14

Query Match 30.5%: Score 30.8; DB 4; Length 1435;
Best Local Similarity 57.1%: Pred. No. 0.11;

Matches 56: Conservative 0; Mismatches 42; Indels 0; Gaps 0;

OY 2 ctcaagttcctgacacatcttggtccctgcgcgcgcctctctctacgcact 61
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OY 62 atccaagattggcattcgacatccgagactgac 99
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Db 677 CTGATGAGCTTGGAATGAGAACGTACCAACAACTTGC 714
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RESULT 3
US-07-866-979-3
Sequence 3, Application US/07866979
Patent No. 5532347
GENERAL INFORMATION:
APPLICANT: Cone, Roger D
APPLICANT: Mountjoy, Kathleen G
TITLE OF INVENTION: Melanocyte Stimulating Hormone Receptor
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Allegritti & Witcoff, Ltd.
STREET: 10 South Wacker Drive, Suite 3000
CITY: Chicago
STATE: Illinois
COUNTRY: USA
ZIP: 60606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/866,979
FILING DATE: 19920410
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: No. 5532347nan, Kevin E
REGISTRATION NUMBER: 35,303
REFERENCE/DOCKET NUMBER: 92,154
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312-715-1000
TELEFAX: 312-715-1234
TELEX: 910-221-5317
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1260 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 15..959
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..14
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 960..1260
US-07-866-979-3

Query Match 29.1%: Score 29.4; DB 1; Length 1260;
Best Local Similarity 58.6%: Pred. No. 0.33;
Matches 51: Conservative 0; Mismatches 36; Indels 0; Gaps 0;

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|||||
Db 363 CTCATGACGTGCTCATCTGTGGCTCCATGCTGTCAGTCTGCTTCGCGCATCATTT 422
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OY 62 atccaagattggcattcgacatccgagactgac 88
- - - - -

Db 423 GCTATAGACCGCTACATCTTCATCTTC 449

```

RESULT      4
US-08-466-906B-3
: Sequence 3, Application US/08466906B
: Patent No. 5849871
:
: GENERAL INFORMATION:
: APPLICANT: Cong, Roger D
: APPLICANT: Mountjoy, Kathleen G
: TITLE OF INVENTION: Melanocyte Stimulating Hormone Receptor
: TITLE OF INVENTION: and Uses
: NUMBER OF SEQUENCES: 8
:
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: McConnell Boehnen Hulbert & Berghoff
: STREET: 300 South Wacker Drive
: CITY: Chicago
:
: STATE: IL
: COUNTRY: USA
: ZIP: 60606
:
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patent Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/466,906B
: FILING DATE: 06-JUN-1995
: CLASSIFICATION: 530
:
: ATTORNEY/AGENT INFORMATION:
: NAME: No. 5849871nan, Kevin E
: REGISTRATION NUMBER: 35,303
: REFERENCE/DOCKET NUMBER: 92,154-H
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 312-913-0001
: TELEFAX: 312-913-0002
:
:
: INFORMATION FOR SEQ ID NO: 3:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1260 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA to mRNA
:
: FEATURE:
: NAME/KEY: 5'UTR
: LOCATION: 1..14
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 15..959
: FEATURE:
: NAME/KEY: 3'UTR
: LOCATION: 960..1260
:
: US-08-466-906B-3

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Query Match	29.1%	Score 29.4	DB 2	Length 1260
Best Local Similarity	58.6%	Pred No. 0.33		
Matches	51	Conservative	0	Mismatches 36
			Indels	0
			Gaps	0
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Db	363	ctctattgacgcgcgtcattctgttcgcctccattgttccactctctctccatcttcatt	422	
Oy	62	atccaaagatttggacatcgcgcacattc	88	
Db	423	gctattagaccgctacattctccattcttc	449	

RESULT 5
US-08-706-281A-3
; Sequence 3, Application US/08706281A

```

? Patent No. 6100048
? GENERAL INFORMATION:
? APPLICANT: Cone, Roger D
? APPLICANT: Fan, Wei
? APPLICANT: Boston, Bruce A
? APPLICANT: Kesteron, Robert A
? APPLICANT: Lu, Dongsi
? APPLICANT: Chen, Wenbiao
? TITLE OF INVENTION: Methods and Reagents for Discovering and
? TITLE OF INVENTION: Using Mammalian Melanocortin Receptor Agonists and Antagonists
? TITLE OF INVENTION: To Modulate Feeding Behavior in Animals
? NUMBER OF SEQUENCES: 19
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: McDonnell Boehnen Hulbert & Berghoff
? STREET: 300 South Wacker Drive
? CITY: Chicago
? STATE: IL
? COUNTRY: USA
? ZIP: 60606
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Floppy disk
? COMPUTER: IBM PC compatible
? OPERATING SYSTEM: PC-DOS/MS-DOS
? SOFTWARE: Patent Release #1.0, Version #1.25
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/706,281A
? FILING DATE: 04-SEP-1996
? CLASSIFICATION: 435
? ATTORNEY/AGENT INFORMATION:
? NAME: No. 6100048nan, Kevin E
? REGISTRATION NUMBER: 35,303
? REFERENCE/DOCKET NUMBER: 96,886
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: 312-913-0001
? TELEFAX: 312-913-0002
? TELEX:
? INFORMATION FOR SEQ ID NO: 3:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 1260 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: single
? TOPOLOGY: linear
? MOLECULE TYPE: cDNA to mRNA
? FEATURE:
? NAME/KEY: 5'UTR
? LOCATION: 1..14
? FEATURE:
? NAME/KEY: CDS
? LOCATION: 15..959
? FEATURE:
? NAME/KEY: 3'UTR
? LOCATION: 960..1260
? US-08-706-281A-3

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Best Local Similarity	58.6%		
Matches	51;	Conservative	0; Mismatches 36; Indels 0; Gaps 0;
Db	363	CTCATTTAGCGCTCATCTTGTGGCTCCATGGTGTCCATGCTCTGTGGTCTGGGCATCATTT	422
Oy	62	atccaagatttgggacatcgacacalttc	88
Db	423	GCTATPAGACGCTCATCTTCATTCGATCTCT	449

RESULT 6
US-09-201-746-3
; Sequence 3, Application US/09201746
; Patent No. 6268221
; GENERAL INFORMATION:

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1  APPLICANT: Cong, Roger D
2  APPLICANT: Mounjoy, Kathleen G
3  TITLE OF INVENTION: Melanocyte Stimulating Hormone Receptor
4  TITLE OF INVENTION: and Uses
5  NUMBER OF SEQUENCES: 8
6  CORRESPONDENCE ADDRESS:
7  ADDRESSEE: McDonnell
8  STREET: 300 South Wacker Drive
9  CITY: Chicago
10 STATE: IL
11 COUNTRY: USA
12 ZIP: 60606
13 COMPUTER READABLE FORM:
14 MEDIUM TYPE: floppy disk
15 COMPUTER: IBM PC compatible
16 OPERATING SYSTEM: PC-DOS/MS-DOS
17 SOFTWARE: Patent Release #1.0, Version #1.25
18 CURRENT APPLICATION DATA:
19 APPLICATION NUMBER: US/09/201,746
20 FILING DATE: 01-DEC-1998
21 CLASSIFICATION: 435
22 ATTORNEY/AGENT INFORMATION:
23 NAME: No. 626821nan, Kevin E
24 REGISTRATION NUMBER: 35,303
25 REFERENCE/DOCKET NUMBER: 92,154-J
26 TELECOMMUNICATION INFORMATION:
27 TELEPHONE: 312-913-0001
28 TELEFAX: 312-913-0002
29 TELEX:
30 INFORMATION FOR SEQ ID NO: 3:
31 SEQUENCE CHARACTERISTICS:
32 LENGTH: 1260 base pairs
33 TYPE: nucleic acid
34 STRANDEDNESS: single
35 TOPOLOGY: linear
36 MOLECULE TYPE: cDNA to mRNA
37 FEATURE:
38 NAME/KEY: 5'UTR
39 LOCATION: 1..14
40 FEATURE:
41 NAME/KEY: CDS
42 LOCATION: 15..959
43 FEATURE:
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45 LOCATION: 960..1260
46
47 US-09-201-746-3
48
49 Query Match 29.1%; Score 29.4; DB 4; Length 1260;
50 Best Local Similarity 58.6%; Pred. No. 0.33;
51 Matches 51; Conservative 0; Mismatches 36; Indels 0; Gaps 0;
52
53 QY 2 ctcaagttccgcacatcttggttccctgctgcccggccctctcctctaccacact 61
54 |||| | ||| |||| | || | || | || | || | || | || | || | || |
55 Db 363 ctcaattgacgtgctcattgtggctccatggtgctccactgtctgcttctgagcatatt 422
56 || | || | || | || | || | || | || | || | || | || | || | || |
57 QY 62 atccaagattcggacatcgacattc 88
58 || | || | || | || | || | || | || | || | || | || | || | || |
59 Db 423 gctatgacgcgctgacatctccatcttc 449
60
61 RESULT 7
62 US-09-097-231-3
63 ; Sequence 3, Application US/09097231
64 ; Patent No. 6278038
65 ;
66 GENERAL INFORMATION:
67 ;
68 APPLICANT: Cong, Roger D
69 ; Chen, Wenbiao
70 ; Low, Malcolm J
71 TITLE OF INVENTION: Mammalian Melanocortin Receptor and Uses
72 ;
73 NUMBER OF SEQUENCES: 22
74 ;
75 CORRESPONDENCE ADDRESS:
76 ;
77 ADDRESSEE: McDonnell Boehnen Hulbert & Berghoff

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1 /
2 / STREET: 300 South Wacker Drive
3 / CITY: Chicago
4 / STATE: Illinois
5 / COUNTRY: USA
6 / ZIP: 60606
7 /
8 / COMPUTER READABLE FORM:
9 / MEDIUM TYPE: Floppy disk
10 / COMPUTER: IBM PC compatible
11 / OPERATING SYSTEM: PC-DOS/MS-DOS
12 / SOFTWARE: Patentin Release #1.0, Version #1.25
13 /
14 / CURRENT APPLICATION DATA:
15 / APPLICATION NUMBER: US/09/097,231
16 / FILING DATE: 12-Jun-1998
17 / CLASSIFICATION: <Unknown>
18 /
19 / ATTORNEY/AGENT INFORMATION:
20 / NAME: No. 6278038aan, Kevin E
21 / REGISTRATION NUMBER: 35,303
22 / REFERENCE/DOCKET NUMBER: 96,886-C
23 / TELECOMMUNICATION INFORMATION:
24 / TELEPHONE: 312-913-0001
25 / TELEFAX: 312-913-0002
26 / TELEX: <Unknown>
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28 / INFORMATION FOR SEQ ID NO: 3:
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30 / SEQUENCE CHARACTERISTICS:
31 / LENGTH: 1260 base pairs
32 / TYPE: nucleic acid
33 / STRANDEDNESS: single
34 / TOPOLOGY: linear
35 / MOLECULE TYPE: cDNA to mRNA
36 /
37 / FEATURE:
38 / NAME/KEY: 5'UTR
39 / LOCATION: 1..14
40 /
41 / FEATURE:
42 / NAME/KEY: CDS
43 / LOCATION: 15..959
44 /
45 / FEATURE:
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47 / LOCATION: 960..1260
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49 / SEQUENCE DESCRIPTION: SEQ ID NO: 3:
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1  COUNTRY: USA
2  ZIP: 07065-0900
3  COMPUTER READABLE FORM:
4  MEDIUM TYPE: Floppy disk
5  COMPUTER: IBM PC compatible
6  OPERATING SYSTEM: PC-DOS/MS-DOS
7  SOFTWARE: Patent in Release #1.0, Version #1.25
8  CURRENT APPLICATION DATA:
9  APPLICATION NUMBER: US/08/759,848
10 PILING DATE:
11 CLASSIFICATION: 800
12 PRIOR APPLICATION DATA:
13 APPLICATION NUMBER: US 08/281,393
14 PILING DATE: 27-Jul-1994
15 ATTORNEY/AGENT INFORMATION:
16 NAME: Wallen III, John W.
17 REGISTRATION NUMBER: 35,403
18 REFERENCE/DOCKET NUMBER: 19234
19 TELECOMMUNICATION INFORMATION:
20 TELEPHONE: (908) 594-3905
21 TELEFAX: (908) 594-4720
22 INFORMATION FOR SEQ ID NO: 2:
23 SEQUENCE CHARACTERISTICS:
24 LENGTH: 1378 base pairs
25 TYPE: nucleic acid
26 STRANDEDNESS: single
27 TOPOLOGY: linear
28 MOLECULE TYPE: DNA (genomic)
29 US-08-759-848-2

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Query Match	27.5%;	Score 27.8;	DB 1;	Length 1378;
Best Local Similarity	65.1%;	Pred. No. 1.2;		
Matches 41;	Conservative 0;	Mismatches 22;	Indels 0;	Gaps 0

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QY	64	cca	66
Db	876	GCA	878

RESULT: 9
 PCT-US95-09383-2
 Sequence 2, Application PC/TUS9509383
 GENERAL INFORMATION:
 APPLICANT: Borkowski, Joseph A.
 APPLICANT: Strader, Catherine D.
 APPLICANT: Hess, John W.
 APPLICANT: Chen, Howard Y.
 APPLICANT: Trumbauer, Myrna E.
 TITLE OF INVENTION: BRADYKININ B2 RECEPTOR MODIFIED
 TITLE OF INVENTION: NON-HUMAN ANIMALS
 NUMBER OF SEQUENCES: 12
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: John W. Wallen III
 STREET: 126 E. Lincoln Avenue
 CITY: Rahway
 STATE: New Jersey
 COUNTRY: USA
 ZIP: 07065-0900
 COMPUTER READABLE FORM:
 MEDIUM TYPE: floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: PCT/US95/09383
 FILING DATE:
 CLASSIFICATION:
 PRIOR APPLICATION DATA:

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1 APPLICATION NUMBER: US 08/281,393
2
3 FILING DATE:
4
5 ATTORNEY/AGENT INFORMATION:
6
7 NAME: Wallen III, John W.
8 REGISTRATION NUMBER: 35,403
9 REFERENCE/DOCKET NUMBER: 19234
10 TELECOMMUNICATION INFORMATION:
11
12 TELEPHONE: (908) 594-3905
13
14 TELEFAX: (908) 594-4720
15
16 INFORMATION FOR SEQ ID NO: 2:
17
18 SEQUENCE CHARACTERISTICS:
19
20 LENGTH: 1378 base pairs
21
22 TYPE: nucleic acid
23 STRANDEDNESS: single
24
25 TOPOLOGY: linear
26
27 MOLECULE TYPE: DNA (genomic)
28
29 PCT-US95-09383-2

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Query Match	27.50;	Score 27.8;	DB 5;	Length 1378;
Best Local Similarity	65.18;	Pred. No. 1.2;		
Matches 41;	Conservative 0;	Mismatches 22;	Indels 0;	Gaps 0;

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QY 64 cca 66
Db 876 GCA 878

RESULT 10
 US-08-938-291A-3/C
 : Sequence 3, Application US/08938291A
 : Patent No. 611673
 :
 : GENERAL INFORMATION:
 : APPLICANT: Lev, Sima
 : APPLICANT: Plozman, Gregory D.
 : APPLICANT: Schlesinger, Joseph
 : TITLE OF INVENTION: RDBG PROTEINS AND RELATED
 : TITLE OF INVENTION: PRODUCTS AND METHODS
 : NUMBER OF SEQUENCES: 11
 : CORRESPONDENCE ADDRESS:
 : ADDRESSEE: Lyon & Lyon
 : STREET: 633 West Fifth Street
 : STREET: Suite 4700
 : CITY: Los Angeles
 : STATE: California
 : COUNTRY: U.S.A.
 : ZIP: 90071-2066
 :
 : COMPUTER READABLE FORM:
 : MEDIUM TYPE: 3.5" Diskette, 1.44 MB
 : MEDIUM TYPE: Storage
 : COMPUTER: IBM Compatible
 : OPERATING SYSTEM: IBM P.C. DOS 5.0
 : SOFTWARE: Fastseq
 :
 : CURRENT APPLICATION DATA:
 : APPLICATION NUMBER: US/08/938,291A
 : FILING DATE: September 26, 1997
 : CLASSIFICATION: 435
 : PRIOR APPLICATION DATA:
 : APPLICATION NUMBER: 60/027,337
 : FILING DATE: October 11, 1996
 : ATTORNEY/AGENT INFORMATION:
 : NAME: Waiburg, Richard J.
 : REESTRICATION NUMBER: 32,327
 : REESTRICATION/DOCKET NUMBER: 228/172
 : TELECOMMUNICATION INFORMATION:
 : TELEPHONE: (213) 489-1600
 : TELEFAX: (213) 955-0440
 : TELEEX: 67-3510
 :
 : INFORMATION FOR SEQ ID NO: 3:


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? Sequence 16, Application US/08479285
? Patent No. 6207815
? GENERAL INFORMATION:
? APPLICANT: MEZES, PETER S
? APPLICANT: GOURLIE, BRIAN B
? APPLICANT: RIXON, MARK W
? APPLICANT: ANDERSON, WH KERR
? APPLICANT: KAPLAN, DONALD A
? APPLICANT: SCHOLOM, JEFFREY
? TITLE OF INVENTION: A NOVEL FAMILY OF HIGH AFFINITY,
? TITLE OF INVENTION: MODIFIED ANTIBODIES FOR CANCER TREATMENT
? NUMBER OF SEQUENCES: 74
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: DUANE C ULMER
? STREET: P O BOX 1967
? CITY: MIDLAND
? STATE: MICHIGAN
? COUNTRY: USA
? ZIP: 48641-1967
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Floppy disk
? COMPUTER: IBM PC compatible
? OPERATING SYSTEM: PC-DOS/MS-DOS
? SOFTWARE: Patentin Release #1.0, Version #1.25
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/479,285
? FILING DATE: 07-JUN-1995
? CLASSIFICATION: 536
? PRIOR APPLICATION DATA:
? APPLICATION NUMBER: US 08/040687
? FILING DATE: 31-MAR-1993
? ATTORNEY/AGENT INFORMATION:
? NAME: ULMER, DUANE C
? REGISTRATION NUMBER: 34,941
? REFERENCE/DOCKET NUMBER: C-37,075C
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: (517) 636-8104
? INFORMATION FOR SEQ ID NO: 16:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 1984 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: double
? TOPOLOGY: linear
? US-08-479-285-16

Query Match 25.3%; Score 25.6; DB 4: Length 1984;
Best Local Similarity 59.7%; Pred. No. 7.9;
Matches 43; Conservative 0; Mismatches 29; Indels 0; Gaps 0;

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DB 897 ctgagatccctgacagatcttttcccaacttctctcagccgctccctcagggcaat 838
OY 62 atccaagattcg 73
DB 837 atccaagattag 826

RESULT 14
US-08-481-337A-1/C
? Sequence 1, Application US/08481337A
? Patent No. 5863738
? GENERAL INFORMATION:
? APPLICANT: TEN DIJKE, Peter
? APPLICANT: HELDIN, Carl-Henrik
? APPLICANT: MIYAZONO, Kohel
? APPLICANT: SAMPATH, Kuber T.
? TITLE OF INVENTION: Morphogenic Protein-Specific Cell
? TITLE OF INVENTION: Surface Receptors and Uses Therefor
? NUMBER OF SEQUENCES: 18
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: Testa, Hurwitz & Thibault
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? STREET: 125 High St.
? CITY: Boston
? STATE: MA
? COUNTRY: USA
? ZIP: 02110
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Floppy disk
? COMPUTER: IBM PC compatible
? OPERATING SYSTEM: PC-DOS/MS-DOS
? SOFTWARE: Patentin Release #1.0, Version #1.30
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/481,337A
? FILING DATE: 02-JUN-1995
? CLASSIFICATION: 435
? ATTORNEY/AGENT INFORMATION:
? NAME: MEYERS, Thomas C.
? REGISTRATION NUMBER: 36,989
? REFERENCE/DOCKET NUMBER: CRP-097CP2
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: (617) 248-7000
? TELEFAX: (617) 248-7100
? INFORMATION FOR SEQ ID NO: 1:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 1509 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: single
? TOPOLOGY: linear
? MOLECULE TYPE: cDNA
? FEATURE:
? NAME/KEY: CDS
? LOCATION: 1..1509
? OTHER INFORMATION: /product="Human ALK1"
? US-08-481-337A-1

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Best Local Similarity 64.4%; Pred. No. 8.5;
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DB 216 caagttccctgcacacacatcttgctccctgcgcgcgcctctctctacaccgact 158

RESULT 15
US-08-696-268B-1/C
? Sequence 1, Application US/08696268B
? Patent No. 5968752
? GENERAL INFORMATION:
? APPLICANT: ICHIO, HIDEORI
? APPLICANT: NISHITO, HIDEKI
? APPLICANT: SAMPATH, KUBER T.
? TITLE OF INVENTION: NOVEL SIGNALING RECEPTOR FOR
? TITLE OF INVENTION: MORPHOGENIC PROTEINS
? NUMBER OF SEQUENCES: 8
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: Testa, Hurwitz & Thibault
? STREET: 125 High St.
? CITY: Boston
? STATE: MA
? COUNTRY: USA
? ZIP: 02110
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Floppy disk
? COMPUTER: IBM PC compatible
? OPERATING SYSTEM: PC-DOS/MS-DOS
? SOFTWARE: Patentin Release #1.0, Version #1.30
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/696,268B
? FILING DATE:
? CLASSIFICATION: 435
? ATTORNEY/AGENT INFORMATION:
? NAME: MEYERS, Thomas C.
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: REGISTRATION NUMBER: 36,989
: REFERENCE/DOCKET NUMBER: CRP-117
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (617) 248-7000
: TELEFAX: (617) 248-7100
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1509 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 1..1509
: OTHER INFORMATION: /product="Human ALK-1"
US-08-696-268B-1
    
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Query Match      25.1% Score 25.4; DB 2; Length 1509;
Best Local Similarity 64.4% Pred No 8.5;
Matches 38: Conservative 0; Mismatches 21; Indels 0; Gaps 0;
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   1111111111111111111111111111111111111111111111111111111
Db 216 CAAGTTCCTCCGACGCCGATGTTCTCTGGGGGTGCTCTCCCTCTCCGACACGACTA 158
    
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Search completed: October 3, 2002, 16:24:54
 Job time: 17134 sec